



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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<p>(21) International Application Number: PCT/US98/06231</p> <p>(22) International Filing Date: 30 March 1998 (30.03.98)</p> <p>(30) Priority Data: 60/042,905 31 March 1997 (31.03.97) US</p> <p>(71) Applicant: YALE UNIVERSITY [US/US]; Suite 401, 246 Church Street, New Haven, CT 06520 (US).</p> <p>(72) Inventor: BREAKER, Ronald, R.; 133 Weatherly Trail, Guilford, CT 06437 (US).</p> <p>(74) Agents: WARBURG, Richard, J. et al.; Lyon &amp; Lyon LLP, Suite 4700, 633 West Fifth Street, Los Angeles, CA 90071-2066 (US).</p>	<p>(81) Designated States: AU, CA, JP, MX, European patent (A.T, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).</p> <p><b>Published</b> <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments,</i></p> <p>(88) Date of publication of the international search report: 15 April 1999 (15.04.99)</p>	
<p>(54) Title: NUCLEIC ACID CATALYSTS</p>		
<p>(57) Abstract</p> <p>Nucleic acid molecules with new motifs having catalytic activity, methods of synthesis, and use thereof are also described.</p>		

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## INTERNATIONAL SEARCH REPORT

Internat. H Application No

PCT/US 98/06231

A. CLASSIFICATION OF SUBJECT MATTER  
IPC 6 C12N9/00 C12N15/09

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  
IPC 6 C12N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	NISHIKAWA, F. ET AL.: "Selection in vitro of trans-acting genomic hepatitis delta virus (HDV) ribozymes" EUR. J. BIOCHEM., vol. 237, 1996, pages 712-718, XP002084352 * fig. 1 *	1,7,13, 17,20,27
X	PERROTTA, A.T. & BEEN, M.D.: "Core sequences and a cleavage site wobble pair required for HDV antigenomic ribozyme self-cleavage" NUCLEIC ACIDS RESEARCH, vol. 24, no. 7, 1996, pages 1314-1321, XP000582452 * fig. 1 *	1,7,13, 17,20,27



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

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"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

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"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"Z" document member of the same patent family

Date of the actual completion of the international search

13 November 1998

Date of mailing of the international search report

03.03.99

Name and mailing address of the ISA

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## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 97 10328 A (RIBOZYME PHARMACEUTICALS) 20 March 1997 * claims *	1
A	--- BREAKER, R.R. & JOYCE, G.F.: "Inventing and improving ribozyme function: ..." TIBTECH, vol. 12, 1994, pages 268-275, XP002084353 *whole disclosure *	1,3,5-34
X	--- WO 95 19433 A (TARGETECH, INC.) 20 July 1995 * claims 15,20,23 *	1,3
X	--- WO 96 03152 A (GEORGETOWN UNIVERSITY). 8 February 1996 * claim 16 *	1,3
X,P	--- TANG, J. & BREAKER, R.R.: "Examination of the catalytic fitness of the hammerhead ribozyme by in vitro selection" RNA, vol. 3, 1997, pages 914-925, XP002084354 * whole disclosure *	1,3,5-34
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# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/IS 98/06231

## Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:  
because they relate to subject matter not required to be searched by this Authority, namely:
  
2. ☐ Claims Nos.:  
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
  
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
  
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
  
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
  
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

1 and 3 completely; 5-34 partially

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

1. Claims: 1 and 3 completely; 5-34 partially

A nucleic acid molecule corresponding to formula I, and related subject-matter

2. Claims: 1 and 3 completely; 5-34 partially

A nucleic acid molecule corresponding to formula II, and related subject-matter.

3. Claims: 35-37

A method of selecting a (structurally undefined, not restricted to formulas I and II) nucleic acid with catalytic activity

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 98/06231

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9710328 A	20-03-1997	AU 6761796 A CA 2226728 A CN 1196091 A EP 0842286 A	01-04-1997 20-03-1997 14-10-1998 20-05-1998
WO 9519433 A	20-07-1995	US 5728518 A AU 1680095 A CA 2180347 A EP 0739415 A JP 9511382 T	17-03-1998 01-08-1995 20-07-1995 30-10-1996 18-11-1998
WO 9603152 A	08-02-1996	US 5646262 A AU 3102495 A CA 2196070 A EP 0772454 A JP 10503367 T	08-07-1997 22-02-1996 08-02-1996 14-05-1997 31-03-1998



## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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<p>(54) Title: <b>NUCLEIC ACID CATALYSTS</b></p> <p>(57) Abstract</p> <p>Nucleic acid molecules with new motifs having catalytic activity, methods of synthesis, and use thereof are also described.</p> <div style="text-align: center;"> <p><b>Cleavage Site</b></p> <p>Target      5' - ..... N N N N U H N N N N N ..... - 3'</p> <p>Ribozyme      3' - ..... N' N' N' N' A<sub>15.1</sub> N' N' N' N' N' ..... - 5'</p> <p>Stem III      Stem I</p> <p>Stem II      Loop II</p> </div> <p>ATP primer      hammerhead</p> <p><b>H3</b></p>		



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DESCRIPTIONNucleic Acid CatalystsBackground of the Invention

5        This invention relates to nucleic acid molecules with catalytic activity and derivatives thereof.

      The following is a brief description of enzymatic nucleic acid molecules. This summary is not meant to be complete but is provided only for understanding of the  
10 invention that follows. This summary is not an admission that all of the work described below is prior art to the claimed invention.

      Enzymatic nucleic acid molecules (ribozymes) are nucleic acid molecules capable of catalyzing one or more  
15 of a variety of reactions, including the ability to repeatedly cleave other separate nucleic acid molecules in a nucleotide base sequence-specific manner. Such enzymatic nucleic acid molecules can be used, for example, to target virtually any RNA transcript (Zaug et al., 324,  
20 Nature 429 1986; Cech, 260 JAMA 3030, 1988; and Jefferies et al., 17 Nucleic Acids Research 1371, 1989).

      Because of their sequence-specificity, trans-cleaving enzymatic nucleic acid molecules show promise as therapeutic agents for human disease (Usman & McSwiggen, 1995  
25 Ann. Rep. Med. Chem. 30, 285-294; Christoffersen and Marr, 1995 J. Med. Chem. 38, 2023-2037). Enzymatic nucleic acid molecules can be designed to cleave specific RNA targets within the background of cellular RNA. Such a cleavage event renders the mRNA non-functional and abrogates  
30 protein expression from that RNA. In this manner, synthesis of a protein associated with a disease state can be selectively inhibited.

There are seven basic varieties of naturally-occurring enzymatic RNAs. Each can catalyze the hydrolysis of RNA phosphodiester bonds in trans (and thus can cleave other RNA molecules) under physiological conditions. In general, enzymatic nucleic acids act by first binding to a target RNA. Such binding occurs through the target binding portion of a enzymatic nucleic acid which is held in close proximity to an enzymatic portion of the molecule that acts to cleave the target RNA. Thus, the enzymatic nucleic acid first recognizes and then binds a target RNA through complementary base-pairing, and once bound to the correct site, acts enzymatically to cut the target RNA. Strategic cleavage of such a target RNA will destroy its ability to direct synthesis of an encoded protein. After an enzymatic nucleic acid has bound and cleaved its RNA target, it is released from that RNA to search for another target and can repeatedly bind and cleave new targets.

In addition, several *in vitro* selection (evolution) strategies (Orgel, 1979, *Proc. R. Soc. London*, B 205, 435) have been used to evolve new nucleic acid catalysts capable of catalyzing a variety of reactions, such as cleavage and ligation of phosphodiester linkages and amide linkages, (Joyce, 1989, *Gene*, 82, 83-87; Beaudry et al., 1992, *Science* 257, 635-641; Joyce, 1992, *Scientific American* 267, 90-97; Breaker et al., 1994, *TIBTECH* 12, 268; Bartel et al., 1993, *Science* 261:1411-1418; Szostak, 1993, *TIBS* 17, 89-93; Kumar et al., 1995, *FASEB J.*, 9, 1183; Breaker, 1996, *Curr. Op. Biotech.*, 7, 442).

The enzymatic nature of a ribozyme is advantageous over other technologies, since the effective concentration of ribozyme necessary to effect a therapeutic treatment is generally lower than that of an antisense oligonucleotide.

This advantage reflects the ability of the ribozyme to act enzymatically. Thus, a single ribozyme (enzymatic nucleic acid) molecule is able to cleave many molecules of target RNA. In addition, the ribozyme is a highly specific inhibitor, with the specificity of inhibition depending not only on the base-pairing mechanism of binding, but also on the mechanism by which the molecule inhibits the expression of the RNA to which it binds. That is, the inhibition is caused by cleavage of the RNA target and so  
10 specificity is defined as the ratio of the rate of cleavage of the targeted RNA over the rate of cleavage of non-targeted RNA. This cleavage mechanism is dependent upon factors additional to those involved in base-pairing. Thus, it is thought that the specificity of action of a  
15 ribozyme is greater than that of antisense oligonucleotide binding the same RNA site.

The development of ribozymes that are optimal for catalytic activity would contribute significantly to any strategy that employs RNA-cleaving ribozymes for the  
20 purpose of regulating gene expression. The hammerhead ribozyme functions with a catalytic rate ( $k_{cat}$ ) of  $\sim 1 \text{ min}^{-1}$  in the presence of saturating (10 mM) concentrations of  $\text{Mg}^{2+}$  cofactor. However, the rate for this ribozyme in  $\text{Mg}^{2+}$  concentrations that are closer to those found inside cells  
25 (0.5 - 2 mM) can be 10- to 100-fold slower. In contrast, the RNase P holoenzyme can catalyze pre-tRNA cleavage with a  $k_{cat}$  of  $\sim 30 \text{ min}^{-1}$  under optimal assay conditions. An artificial 'RNA ligase' ribozyme has been shown to catalyze the corresponding self-modification reaction with  
30 a rate of  $\sim 100 \text{ min}^{-1}$ . In addition, it is known that certain modified hammerhead ribozymes that have substrate binding arms made of DNA catalyze RNA cleavage with multiple turnover rates that approach  $100 \text{ min}^{-1}$ . Finally, replacement

of a specific residue within the catalytic core of the hammerhead with certain nucleotide analogues gives modified ribozymes that show as much as a 10-fold improvement in catalytic rate. These findings demonstrate  
5 that ribozymes can promote chemical transformations with catalytic rates that are significantly greater than those displayed in vitro by most natural self-cleaving ribozymes. It is then possible that the structures of certain self-cleaving ribozymes may not be optimized to  
10 give maximal catalytic activity, or that entirely new RNA motifs could be made that display significantly faster rates for RNA phosphoester cleavage.

An extensive array of site-directed mutagenesis studies have been conducted with the hammerhead ribozyme  
15 to probe relationships between nucleotide sequence and catalytic activity. These systematic studies have made clear that most nucleotides in the conserved core of the hammerhead ribozyme cannot be mutated without significant loss of catalytic activity. In contrast, a combinatorial  
20 strategy that simultaneously screens a large pool of mutagenized ribozymes for RNAs that retain catalytic activity could be used more efficiently to define immutable sequences and to identify new ribozyme variants. Although similar in vitro selection experiments have been  
25 conducted with the hammerhead ribozyme (Nakamaye & Eckstein, 1994, *Biochemistry* 33, 1271; Long & Uhlenbeck, 1994, *Proc. Natl. Acad. Sci.*, 91, 6977; Ishizaka et al., 1995, *BBRC* 214, 403), none of these efforts have successfully screened full-sized hammerhead ribozymes for  
30 all possible combinations of sequence variants that encompass the entire catalytic core.

The references cited above are distinct from the presently claimed invention since they do not disclose

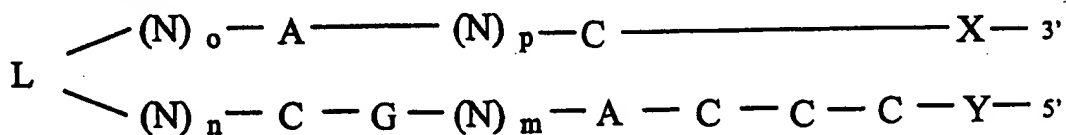
and/or contemplate the enzymatic nucleic acid molecules of the instant invention.

### Summary of the Invention

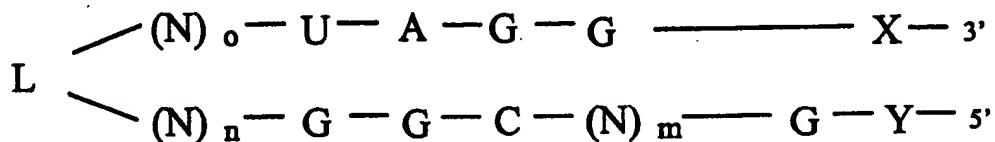
This invention relates to novel nucleic acid molecules with catalytic activity, which are particularly useful for cleavage of RNA or DNA. The nucleic acid catalysts of the instant invention are distinct from other nucleic acid catalysts known in the art. The nucleic acid catalysts of the instant invention do not share sequence homology with other known ribozymes. Specifically, nucleic acid catalysts of the instant invention are capable of catalyzing an intermolecular or intramolecular endonuclease reaction.

In a preferred embodiment, the invention features a nucleic acid molecule with catalytic activity having one of the formulae I-V:

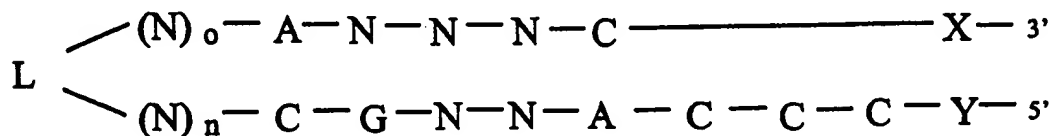
#### Formula I

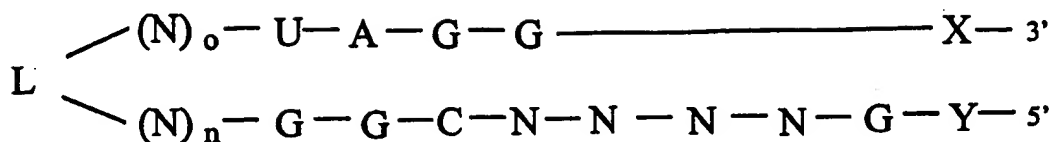
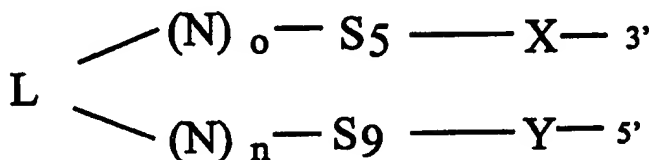


#### 20 Formula II



#### Formula III



Formula IVFormula V

5

In each of the above formula, N represents independently a nucleotide or a non-nucleotide linker, which may be same or different; X and Y are independently oligonucleotides of length sufficient to stably interact (e.g., by forming hydrogen bonds with complementary nucleotides in the target) with a target nucleic acid molecule (the target can be an RNA, DNA or RNA/DNA mixed polymers); o and n are integers greater than or equal to 1 and preferably less than about 100, wherein if (N)<sub>o</sub> and (N)<sub>n</sub> are nucleotides, (N)<sub>o</sub> and (N)<sub>n</sub> are optionally able to interact by hydrogen bond interaction; p and m are independently one of the integers 0, 1, 2, 3, 4 or 5; L is a linker which may be present or absent (i.e., the molecule is assembled from two separate molecules), but when present, is a nucleotide and/or a non-nucleotide linker, which may be a single-stranded and/or double-stranded region; and \_\_\_\_\_ represents a chemical linkage (e.g. a phosphate ester linkage, amide linkage or others known in the art). A, C, U and G represent adenosine, cytidine, uridine and guanosine nucleotides, respectively. The nucleotides in the each of the formula I-V are unmodified or modified at the sugar, base, and/or phosphate as known in the art. S<sub>5</sub> is an oligonucleotide

containing a sequence selected from the group consisting of 5'-AUGUC-3', 5'-ACGUC-3', 5'-ACGGC-3', 5'-ACCUC-3', 5'-AAGGC-3', 5'-AUGGC-3', 5'-AUGCC-3', 5'-ACUCC-3', 5'-AUGAC-3', 5'-ACGAC-3', 5'-UUAGG-3', and 5'-CUAGG-3'; S<sub>9</sub> is an  
5 oligonucleotide containing a sequence selected from the group consisting of 5'-CCCAGUGCC-3', 5'-CCCAGUGCA-3', 5'-CCCAUGCA-3', 5'-CCCAUGCC-3', 5'-CCCAUGCU-3', 5'-CCCAUAGCA-3', 5'-CCCAACGCA-3', 5'-CCCACCGCA-3', 5'-CCCAGAGCA-3', 5'-CCCAACGCU-3', 5'-CCCAUUGCA-3',  
10 GUAGACGGA-3', and 5'-GGUUUCGGG-3'. The nucleotides in the formula V are unmodified or modified at the sugar, base, and/or phosphate portions as known in the art.

In a preferred embodiment, the invention features nucleic acid molecules of any of Formulae I-IV further  
15 comprising a cytidine residue immediately 3' of (N)<sub>n</sub>.

In yet another embodiment, the nucleotide linker (L) is a nucleic acid aptamer, such as an ATP aptamer, HIV Rev aptamer (RRE), HIV Tat aptamer (TAR) and others (for a review see Gold et al., 1995, *Annu. Rev. Biochem.*, 64,  
20 763; and Szostak & Ellington, 1993, in *The RNA World*, ed. Gesteland and Atkins, pp 511, CSH Laboratory Press). A "nucleic acid aptamer" as used herein is meant to indicate nucleic acid sequence capable of interacting with a  
25 molecule, including but not limited to a resin, metabolites, nucleosides, nucleotides, drugs, toxins, transition state analogs, peptides, lipids, proteins, aminoacids, nucleic acid molecules, hormones, carbohydrates, receptors, cells, viruses, bacteria and  
30 others.

In yet another embodiment, the non-nucleotide linker (L) is as defined herein.



The term "nucleotide" is used as recognized in the art to include natural bases, and modified bases well known in the art. Such bases are generally located at the 1' position of a sugar moiety. Nucleotide generally comprise a base, sugar and a phosphate group. The nucleotides can be unmodified or modified at the sugar, phosphate and/or base moiety, (see for example, Usman and McSwiggen, *supra*; Eckstein et al., International PCT Publication No. WO 92/07065; Usman et al., International PCT Publication No. WO 93/15187; all hereby incorporated by reference herein). There are several examples of modified nucleic acid bases known in the art and has recently been summarized by Limbach et al., 1994, *Nucleic Acids Res.* 22, 2183. Some of the non-limiting examples of base modifications that can be introduced into enzymatic nucleic acids without significantly effecting their catalytic activity include, inosine, purine, pyridin-4-one, pyridin-2-one, phenyl, pseudouracil, 2, 4, 6-trimethoxy benzene, 3-methyluracil, dihydrouridine, naphthyl, aminophenyl, 5-alkylcytidines (e.g., 5-methylcytidine), 5-alkyluridines (e.g., ribothymidine), 5-halouridine (e.g., 5-bromouridine) or 6-azapyrimidines or 6-alkylpyrimidines (e.g. 6-methyluridine) and others (Burgin et al., 1996, *Biochemistry*, 35, 14090). By "modified bases" in this aspect is meant nucleotide bases other than adenine, guanine, cytosine and uracil at 1' position or their equivalents; such bases may be used within the catalytic core of the enzyme and/or in the substrate-binding regions.

In particular, the invention features modified ribozymes having a base substitution selected from pyridin-4-one, pyridin-2-one, phenyl, pseudouracil, 2, 4,

6-trimethoxy benzene, 3-methyluracil, dihydrouracil, naphthyl, 6-methyl-uracil and aminophenyl.

There are several examples in the art describing sugar and phosphate modifications that can be introduced  
5 into enzymatic nucleic acid molecules without significantly effecting catalysis and with significant enhancement in their nuclease stability and efficacy. Ribozymes are modified to enhance stability and/or enhance catalytic activity by modification with nuclease resistant  
10 groups, for example, 2'-amino, 2'-C-allyl, 2'-fluoro, 2'-O-methyl, 2'-H, nucleotide base modifications (for a review see Usman and Cedergren, 1992 *TIBS* 17, 34; Usman et al., 1994 *Nucleic Acids Symp. Ser.* 31, 163; Burgin et al., 1996 *Biochemistry* 35, 14090). Sugar modification of  
15 enzymatic nucleic acid molecules have been extensively described in the art (see Eckstein et al., *International Publication* PCT No. WO 92/07065; Perrault et al. *Nature* 1990, 344, 565-568; Pieken et al. *Science* 1991, 253, 314-317; Usman and Cedergren, *Trends in Biochem. Sci.* 1992,  
20 17, 334-339; Usman et al. *International Publication* PCT No. WO 93/15187; Sproat, *US Patent* No. 5,334,711 and Beigelman et al., 1995 *J. Biol. Chem.* 270, 25702).

Such publications describe general methods and strategies to determine the location of incorporation of  
25 sugar, base and/or phosphate modifications and the like into ribozymes without inhibiting catalysis, and are incorporated by reference herein. In view of such teachings, similar modifications can be used as described herein to modify the nucleic acid catalysts of the  
30 instant invention.

In yet another embodiment, the non-nucleotide linker (L) is as defined herein. The term "non-nucleotide" as used herein include either abasic nucleotide, polyether,

polyamine, polyamide, peptide, carbohydrate, lipid, or polyhydrocarbon compounds. Specific examples include those described by Seela and Kaiser, *Nucleic Acids Res.* 1990, 18:6353 and *Nucleic Acids Res.* 1987, 15:3113; Cload  
5 and Schepartz, *J. Am. Chem. Soc.* 1991, 113:6324; Richardson and Schepartz, *J. Am. Chem. Soc.* 1991, 113:5109; Ma et al., *Nucleic Acids Res.* 1993, 21:2585 and *Biochemistry* 1993, 32:1751; Durand et al., *Nucleic Acids Res.* 1990, 18:6353; McCurdy et al., *Nucleosides &*  
10 *Nucleotides* 1991, 10:287; Jsckke et al., *Tetrahedron Lett.* 1993, 34:301; Ono et al., *Biochemistry* 1991, 30:9914; Arnold et al., International Publication No. WO 89/02439; Usman et al., International Publication No. WO 95/06731; Dudycz et al., International Publication No. WO 95/11910  
15 and Ferentz and Verdine, *J. Am. Chem. Soc.* 1991, 113:4000, all hereby incorporated by reference herein. Thus, in a preferred embodiment, the invention features an enzymatic nucleic acid molecule having one or more non-nucleotide moieties, and having enzymatic activity to cleave an RNA  
20 or DNA molecule. By the term "non-nucleotide" is meant any group or compound which can be incorporated into a nucleic acid chain in the place of one or more nucleotide units, including either sugar and/or phosphate substitutions, and allows the remaining bases to exhibit  
25 their enzymatic activity. The group or compound is abasic in that it does not contain a commonly recognized nucleotide base, such as adenosine, guanine, cytosine, uracil or thymine. The terms "abasic" or "abasic nucleotide" as used herein encompass sugar moieties  
30 lacking a base or having other chemical groups in place of base at the 1' position.

In preferred embodiments, the enzymatic nucleic acid includes one or more stretches of RNA, which provide the

enzymatic activity of the molecule, linked to the non-nucleotide moiety. The necessary RNA components are known in the art, see, e.g., Usman, *supra*. By RNA is meant a molecule comprising at least one ribonucleotide residue.

5 As the term is used in this application, non-nucleotide-containing enzymatic nucleic acid means a nucleic acid molecule that contains at least one non-nucleotide component which replaces a portion of a ribozyme, e.g., but not limited to, a double-stranded  
10 stem, a single-stranded "catalytic core" sequence, a single-stranded loop or a single-stranded recognition sequence. These molecules are able to cleave (preferably, repeatedly cleave) separate RNA or DNA molecules in a nucleotide base sequence specific manner. Such molecules  
15 can also act to cleave intramolecularly if that is desired. Such enzymatic molecules can be targeted to virtually any RNA transcript.

By the phrase "nucleic acid catalyst" is meant a nucleic acid molecule capable of catalyzing (altering the  
20 velocity and/or rate of) a variety of reactions including the ability to repeatedly cleave other separate nucleic acid molecules (endonuclease activity) in a nucleotide base sequence-specific manner. Such a molecule with endonuclease activity may have complementarity in a  
25 substrate binding region (e.g. X and Y in formulae I-V) to a specified gene target, and also has an enzymatic activity that specifically cleaves RNA or DNA in that target. That is, the nucleic acid molecule with endonuclease activity is able to intramolecularly or  
30 intermolecularly cleave RNA or DNA and thereby inactivate a target RNA or DNA molecule. This complementarity functions to allow sufficient hybridization of the enzymatic RNA molecule to the target RNA or DNA to allow the cleavage to occur. 100% complementarity is preferred,

but complementarity as low as 50-75% may also be useful in this invention. The nucleic acids may be modified at the base, sugar, and/or phosphate groups. The term enzymatic nucleic acid is used interchangeably with phrases such as  
5 ribozymes, catalytic RNA, enzymatic RNA, catalytic DNA, nucleozyme, DNAzyme, RNA enzyme, endoribonuclease, endonuclease, minizyme, leadzyme, oligozyme or DNA enzyme. All of these terminologies describe nucleic acid molecules with enzymatic activity.

10 By "nucleic acid molecule" as used herein is meant a molecule comprising nucleotides. The nucleic acid can be composed of modified or unmodified nucleotides or non-nucleotides or various mixtures and combinations thereof.

By "complementarity" is meant a nucleic acid that can  
15 form hydrogen bond(s) with other RNA sequence by either traditional Watson-Crick or other non-traditional types (for example, Hoogsteen type) of base-paired interactions.

By "oligonucleotide" as used herein, is meant a molecule comprising two or more nucleotides.

20 The specific enzymatic nucleic acid molecules described in the instant application are not limiting in the invention and those skilled in the art will recognize that all that is important in an enzymatic nucleic acid molecule of this invention is that it has a specific  
25 substrate binding site (e.g., X and/or Y of Formulae 1-V above) which is complementary to one or more of the target nucleic acid regions, and that it have nucleotide sequences within or surrounding that substrate binding site which impart a nucleic acid cleaving activity to the  
30 molecule.

The invention provides a method for producing a class of enzymatic cleaving agents which exhibit a high degree of specificity for the nucleic acid sequence of a desired target. The enzymatic nucleic acid molecule is preferably

targeted to a highly conserved sequence region of a target such that specific diagnosis and/or treatment of a disease or condition can be provided with a single enzymatic nucleic acid. Such enzymatic nucleic acid molecules can  
5 be delivered exogenously to specific cells as required. In the preferred hammerhead motif the small size (less than 60 nucleotides, preferably between 30-40 nucleotides in length) of the molecule allows the cost of treatment to be reduced.

10 Therapeutic ribozymes must remain stable within cells until translation of the target RNA has been inhibited long enough to reduce the levels of the undesirable protein. This period of time varies between hours to days depending upon the disease state. Clearly, ribozymes must  
15 be resistant to nucleases in order to function as effective intracellular therapeutic agents. Improvements in the chemical synthesis of RNA (Wincott et al., 1995 *Nucleic Acids Res.* 23, 2677; incorporated by reference herein) have expanded the ability to modify ribozymes to  
20 enhance their nuclease stability.

By "enzymatic portion" is meant that part of the ribozyme essential for cleavage of an RNA substrate.

By "substrate binding arm" is meant that portion of a ribozyme which is complementary to (i.e., able to base-  
25 pair with) a portion of its substrate. Generally, such complementarity is 100%, but can be less if desired. For example, as few as 10 bases out of 14 may be base-paired. Such arms are shown generally in Figure 1A and as X and/or Y in Formulae I-V. That is, these arms contain sequences  
30 within a ribozyme which are intended to bring ribozyme and target RNA together through complementary base-pairing interactions.

In a preferred embodiment, the invention provides a method for producing a class of enzymatic cleaving agents which exhibit a high degree of specificity for the nucleic acid of a desired target. Such enzymatic nucleic acid molecules can be delivered exogenously to specific cells as required. Alternatively, the ribozymes can be expressed from DNA/RNA vectors that are delivered to specific cells.

The enzymatic nucleic acid molecules of the instant invention can also be expressed within cells from eukaryotic promoters (e.g., Izant and Weintraub, 1985 *Science* 229, 345; McGarry and Lindquist, 1986 *Proc. Natl. Acad. Sci. USA* 83, 399; SullengerScanlon et al., 1991, *Proc. Natl. Acad. Sci. USA*, 88, 10591-5; Kashani-Sabet et al., 1992 *Antisense Res. Dev.*, 2, 3-15; Dropulic et al., 1992 *J. Virol*, 66, 1432-41; Weerasinghe et al., 1991 *J. Virol*, 65, 5531-4; Ojwang et al., 1992 *Proc. Natl. Acad. Sci. USA* 89, 10802-6; Chen et al., 1992 *Nucleic Acids Res.*, 20, 4581-9; Sarver et al., 1990 *Science* 247, 1222-1225; Thompson et al., 1995 *Nucleic Acids Res.* 23, 2259). Those skilled in the art realize that any nucleic acid can be expressed in eukaryotic cells from the appropriate DNA/RNA vector. The activity of such nucleic acids can be augmented by their release from the primary transcript by a ribozyme (Draper et al., PCT WO93/23569, and Sullivan et al., PCT WO94/02595; Ohkawa et al., 1992 *Nucleic Acids Symp. Ser.*, 27, 15-6; Taira et al., 1991, *Nucleic Acids Res.*, 19, 5125-30; Ventura et al., 1993 *Nucleic Acids Res.*, 21, 3249-55; Chowrira et al., 1994 *J. Biol. Chem.* 269, 25856; hereby incorporated in their totality by reference herein).

By "consists essentially of" is meant that the active ribozyme contains an enzymatic center or core equivalent

to those in the examples, and binding arms able to bind target nucleic acid molecules such that cleavage at the target site occurs. Other sequences may be present which do not interfere with such cleavage.

5 Thus, in one aspect, the invention features ribozymes that inhibit gene expression and/or cell proliferation. These chemically or enzymatically synthesized nucleic acid molecules contain substrate binding domains that bind to accessible regions of specific target nucleic acid  
10 molecules. The nucleic acid molecules also contain domains that catalyze the cleavage of target. Upon binding, the enzymatic nucleic acid molecules cleave the target molecules, preventing for example, translation and protein accumulation. In the absence of the expression of  
15 the target gene, cell proliferation, for example, is inhibited.

In a preferred embodiment, the enzymatic nucleic acid molecules are added directly, or can be complexed with cationic lipids, packaged within liposomes, or otherwise  
20 delivered to smooth muscle cells. The RNA or RNA complexes can be locally administered to relevant tissues through the use of a catheter, infusion pump or stent, with or without their incorporation in biopolymers. Using the methods described herein, other enzymatic nucleic acid  
25 molecules that cleave target nucleic acid may be derived and used as described above. Specific examples of nucleic acid catalysts of the instant invention are provided below in the Tables and figures.

Sullivan, et al., *supra*, describes the general  
30 methods for delivery of enzymatic RNA molecules. Ribozymes may be administered to cells by a variety of methods known to those familiar to the art, including, but not restricted to, encapsulation in liposomes, by iontophoresis, or by incorporation into other vehicles, such as



hydrogels, cyclodextrins, biodegradable nanocapsules, and bioadhesive microspheres. For some indications, ribozymes may be directly delivered *ex vivo* to cells or tissues with or without the aforementioned vehicles. Alternatively, the RNA/vehicle combination is locally delivered by direct injection or by use of a catheter, infusion pump or stent. Other routes of delivery include, but are not limited to, intravascular, intramuscular, subcutaneous or joint injection, aerosol inhalation, oral (tablet or pill form), topical, systemic, ocular, intraperitoneal and/or intrathecal delivery. More detailed descriptions of ribozyme delivery and administration are provided in Sullivan *et al.*, *supra* and Draper *et al.*, *supra* which have been incorporated by reference herein.

In another aspect of the invention, enzymatic nucleic acid molecules that cleave target molecules are expressed from transcription units inserted into DNA or RNA vectors. The recombinant vectors are preferably DNA plasmids or viral vectors. Ribozyme expressing viral vectors could be constructed based on, but not limited to, adeno-associated virus, retrovirus, adenovirus, or alphavirus. Preferably, the recombinant vectors capable of expressing the ribozymes are delivered as described above, and persist in target cells. Alternatively, viral vectors may be used that provide for transient expression of ribozymes. Such vectors might be repeatedly administered as necessary. Once expressed, the ribozymes cleave the target mRNA. Delivery of ribozyme expressing vectors could be systemic, such as by intravenous or intramuscular administration, by administration to target cells *ex-planted* from the patient followed by reintroduction into the patient, or by any other means that would allow for introduction into the desired target cell (for a review see Couture and Stinchcomb, 1996, *TIG.*, 12, 510).

In a preferred embodiment, an expression vector comprising nucleic acid sequence encoding at least one of the nucleic acid catalyst of the instant invention is disclosed. The nucleic acid sequence encoding the nucleic acid catalyst of the instant invention is operable linked in a manner which allows expression of that nucleic acid molecule.

In one embodiment, the expression vector comprises: a transcription initiation region (e.g., eukaryotic pol I, II or III initiation region); b) a transcription termination region (e.g., eukaryotic pol I, II or III termination region); c) a gene encoding at least one of the nucleic acid catalyst of the instant invention; and wherein said gene is operably linked to said initiation region and said termination region, in a manner which allows expression and/or delivery of said nucleic acid molecule. The vector may optionally include an open reading frame (ORF) for a protein operably linked on the 5' side or the 3'-side of the gene encoding the nucleic acid catalyst of the invention; and/or an intron (intervening sequences).

By "patient" is meant an organism which is a donor or recipient of explanted cells or the cells themselves. "Patient" also refers to an organism to which enzymatic nucleic acid molecules can be administered. Preferably, a patient is a mammal or mammalian cells. More preferably, a patient is a human or human cells.

By "vectors" is meant any nucleic acid- and/or viral-based technique used to deliver a desired nucleic acid.

Another means of accumulating high concentrations of a ribozyme(s) within cells is to incorporate the ribozyme-encoding sequences into a DNA or RNA expression vector. Transcription of the ribozyme sequences are driven from a promoter for eukaryotic RNA polymerase I (pol I), RNA

polymerase II (pol II), or RNA polymerase III (pol III). Transcripts from pol II or pol III promoters will be expressed at high levels in all cells; the levels of a given pol II promoter in a given cell type will depend on the nature of the gene regulatory sequences (enhancers, silencers, etc.) present nearby. Prokaryotic RNA polymerase promoters are also used, providing that the prokaryotic RNA polymerase enzyme is expressed in the appropriate cells (Elroy-Stein and Moss, 1990 *Proc. Natl. Acad. Sci. U S A*, 87, 6743-7; Gao and Huang 1993 *Nucleic Acids Res.*, 21, 2867-72; Lieber et al., 1993 *Methods Enzymol.*, 217, 47-66; Zhou et al., 1990 *Mol. Cell. Biol.*, 10, 4529-37). Several investigators have demonstrated that ribozymes expressed from such promoters can function in mammalian cells (e.g. Kashani-Sabet et al., 1992 *Antisense Res. Dev.*, 2, 3-15; Ojwang et al., 1992 *Proc. Natl. Acad. Sci. U S A*, 89, 10802-6; Chen et al., 1992 *Nucleic Acids Res.*, 20, 4581-9; Yu et al., 1993 *Proc. Natl. Acad. Sci. U S A*, 90, 6340-4; L'Huillier et al., 1992 *EMBO J.* 11, 4411-8; Lisziewicz et al., 1993 *Proc. Natl. Acad. Sci. U. S. A.*, 90, 8000-4; Thompson et al., 1995 *Nucleic Acids Res.* 23, 2259; Sullenger & Cech, 1993, *Science*, 262, 1566). The above ribozyme transcription units can be incorporated into a variety of vectors for introduction into mammalian cells, including but not restricted to, plasmid DNA vectors, viral DNA vectors (such as adenovirus or adeno-associated virus vectors), or viral RNA vectors (such as retroviral or alphavirus vectors) (for a review see Couture and Stinchcomb, 1996, *supra*).

In a preferred embodiment, the invention features a method of synthesis of enzymatic nucleic acid molecules of instant invention which follows the procedure for normal

chemical synthesis of RNA as described in Usman et al., 1987 *J. Am. Chem. Soc.*, 109, 7845; Scaringe et al., 1990 *Nucleic Acids Res.*, 18, 5433; and Wincott et al., 1995 *Nucleic Acids Res.* 23, 2677-2684 and makes use of common  
5 nucleic acid protecting and coupling groups, such as dimethoxytrityl at the 5'-end, and phosphoramidites at the 3'-end. Small scale synthesis were conducted on a 394 Applied Biosystems, Inc. synthesizer using a modified 2.5  $\mu$ mol scale protocol with a 5 min coupling step for  
10 alkylsilyl protected nucleotides and 2.5 min coupling step for 2'-O-methylated nucleotides. Table II outlines the amounts, and the contact times, of the reagents used in the synthesis cycle. A 6.5-fold excess (163  $\mu$ L of 0.1 M = 16.3  $\mu$ mol) of phosphoramidite and a 24-fold excess of S-  
15 ethyl tetrazole (238  $\mu$ L of 0.25 M = 59.5  $\mu$ mol) relative to polymer-bound 5'-hydroxyl is used in each coupling cycle. Average coupling yields on the 394 Applied Biosystems, Inc. synthesizer, determined by colorimetric quantitation of the trityl fractions, is 97.5-99%. Other  
20 oligonucleotide synthesis reagents for the 394 Applied Biosystems, Inc. synthesizer : detritylation solution was 2% TCA in methylene chloride (ABI); capping was performed with 16% N-methyl imidazole in THF (ABI) and 10% acetic anhydride/10% 2,6-lutidine in THF (ABI); oxidation  
25 solution was 16.9 mM I<sub>2</sub>, 49 mM pyridine, 9% water in THF (Millipore). B & J Synthesis Grade acetonitrile is used directly from the reagent bottle. S-Ethyl tetrazole solution (0.25 M in acetonitrile) is made up from the solid obtained from American International Chemical, Inc.  
30 In a preferred embodiment, deprotection of the chemically synthesized nucleic acid catalysts of the invention is performed as follows. The polymer-bound oligoribonucleotide, trityl-off, is transferred from the

synthesis column to a 4mL glass screw top vial and suspended in a solution of methylamine (MA) at 65 °C for 10 min. After cooling to -20 °C, the supernatant is removed from the polymer support. The support is washed 5 three times with 1.0 mL of EtOH:MeCN:H<sub>2</sub>O/3:1:1, vortexed and the supernatant is then added to the first supernatant. The combined supernatants, containing the oligoribonucleotide, are dried to a white powder.

The base-deprotected oligoribonucleotide is 10 resuspended in anhydrous TEA·HF/NMP solution (250 µL of a solution of 1.5mL N-methylpyrrolidinone, 750 µL TEA and 1.0 mL TEA·3HF to provide a 1.4M HF concentration) and heated to 65°C for 1.5 h. The resulting, fully deprotected, oligomer is quenched with 50 mM TEAB (9 mL) 15 prior to anion exchange desalting.

For anion exchange desalting of the deprotected oligomer, the TEAB solution is loaded on to a Qiagen 500® anion exchange cartridge (Qiagen Inc.) that is prewashed with 50 mM TEAB (10 mL). After washing the loaded 20 cartridge with 50 mM TEAB (10 mL), the RNA is eluted with 2 M TEAB (10 mL) and dried down to a white powder. The average stepwise coupling yields are generally >98% (Wincott et al., 1995 *Nucleic Acids Res.* 23, 2677-2684).

Ribozymes of the instant invention are also 25 synthesized from DNA templates using bacteriophage T7 RNA polymerase (Milligan and Uhlenbeck, 1989, *Methods Enzymol.* 180, 51).

Ribozymes are purified by gel electrophoresis using general methods or are purified by high pressure liquid 30 chromatography (HPLC; See Wincott et al., *supra*) the totality of which is hereby incorporated herein by reference) and are resuspended in water.

In another preferred embodiment, catalytic activity of the molecules described in the instant invention can be optimized as described by Draper et al., *supra*. The details will not be repeated here, but include altering the length of the ribozyme binding arms, or chemically synthesizing ribozymes with modifications (base, sugar and/or phosphate) that prevent their degradation by serum ribonucleases and/or enhance their enzymatic activity (see e.g., Eckstein et al., International Publication No. WO 92/07065; Perrault et al., 1990 *Nature* 344, 565; Pieken et al., 1991 *Science* 253, 314; Usman and Cedergren, 1992 *Trends in Biochem. Sci.* 17, 334; Usman et al., International Publication No. WO 93/15187; and Rossi et al., International Publication No. WO 91/03162; Sproat, US Patent No. 5,334,711; and Burgin et al., *supra*; all of these describe various chemical modifications that can be made to the base, phosphate and/or sugar moieties of enzymatic RNA molecules). Modifications which enhance their efficacy in cells, and removal of bases from stem loop structures to shorten RNA synthesis times and reduce chemical requirements are desired. (All these publications are hereby incorporated by reference herein.).

By "enhanced enzymatic activity" is meant to include activity measured in cells and/or *in vivo* where the activity is a reflection of both catalytic activity and ribozyme stability. In this invention, the product of these properties is increased or not significantly (less than 10 fold) decreased *in vivo* compared to an all RNA ribozyme.

In yet another preferred embodiment, nucleic acid catalysts having chemical modifications which maintain or enhance enzymatic activity is provided. Such nucleic acid

is also generally more resistant to nucleases than unmodified nucleic acid. Thus, in a cell and/or *in vivo* the activity may not be significantly lowered. As exemplified herein such ribozymes are useful in a cell and/or *in vivo* even if activity over all is reduced 10 fold (Burgin et al., 1996, *Biochemistry*, 35, 14090). Such ribozymes herein are said to "maintain" the enzymatic activity on all RNA ribozyme.

Other features and advantages of the invention will be apparent from the following description of the preferred embodiments thereof, and from the claims.

#### Description of the Preferred Embodiments

The drawings will first briefly be described.

#### Drawings:

Figure 1 A) is a diagrammatic representation of the hammerhead ribozyme domain known in the art. Stem II can be  $\geq 2$  base-pair long. Each N is independently any base or non-nucleotide as used herein. B) is a schematic representation of an ATP-dependent ribozyme (H3).

Figure 2 A-C are diagrammatic representations of self-cleaving ribozyme constructs. The arrow head indicates the site of cleavage. N5 and N9 represent the regions of randomization. N9, nine nucleotides in the region from position 16-24 were randomized. N5, five nucleotides in the region from position 59-63 were randomized.

Figure 3 is a schematic representation of a non-limiting *in vitro* selection strategy (allosteric delay) used to evolve nucleic acid catalysts. RT-PCR indicates reverse transcription (RT) and polymerase chain reaction (PCR). This step involves the conversion of RNA into complementary DNA using reverse transcriptase enzyme

followed by PCR amplification to generate a double stranded DNA template for further rounds of selection.

Figure 4 shows a comparison of the yields of RNA self-cleavage from AD-H2 and the generation 4 (G4) and generation 6 (G6) RNA pools in buffer B. Shaded and open bars depict the fraction of precursor cleaved in the presence and absence of ATP, respectively.

Figure 5 Sequences of individual RNAs that represent two new classes of self-cleaving ribozymes. A: Class I ribozymes, that were cloned from populations G6 (generation 6), G7-ATP (generation 7, without ATP), or G9-low (generation 9 in low magnesium concentration) as indicated, are defined by the presence of similar sequences in the regions that correspond to the N<sub>9</sub> and N<sub>5</sub> random-sequence domains of AD-H2. Each member of class I has also acquired a single G to C mutation at nucleotide 28 of the mutagenized AD-H2 construct. B: Class II ribozymes were represented only once (v1) in 34 sequences analyzed from G6. This individual has acquired two additional mutations at nucleotides 35 (A to G) and 37 (U to A) in the aptamer domain of mutagenized AD-H2. An additional class II ribozyme (v2) was isolated from the G9-low population. This variant has the same two mutations as class II v1, but has also acquired a C to A mutation at position 51, and a G deletion at position 32, each relative to the mutagenized AD-H2 construct.

Figure 6 Sequence conservation in the N<sub>9</sub> and N<sub>5</sub> domains among 24 individual class I ribozymes. The frequency of sequence variation compared to clone v2 is plotted as stacked bars, where component bars indicate the contribution to sequence variation for individual nucleotides.



Figure 7 Sequence and possible secondary structure of the constructs 'v2 trans' used to assess the catalytic activity of separate enzyme and substrate domains of a class I ribozyme. Nucleotides in the substrate-binding arms have been altered to complement the corresponding substrate RNA. Encircled nucleotides match those nucleotides that are characteristic of the class I ribozyme 'v2' (Figure 5A). Arrowhead identifies the new site of cleavage.

## 10 Examples

The following are non-limiting examples showing the selection, isolation, synthesis and activity of enzymatic nucleic acids of the instant invention.

The development of nucleic acid catalysts that are optimal for catalytic activity would contribute significantly to any strategy that employs nucleic acid cleaving ribozymes for the purpose of regulating gene expression. The hammerhead ribozyme functions with a catalytic rate ( $k_{cat}$ ) of  $\sim 1 \text{ min}^{-1}$  in the presence of saturating (10 mM) concentrations of  $\text{Mg}^{2+}$  cofactor. However, the rate for this ribozyme in  $\text{Mg}^{2+}$  concentrations that are closer to those found inside cells (0.5 - 2 mM) may be 10- to 100-fold slower. In contrast, the RNase P holoenzyme is believed to catalyze pre-tRNA cleavage with a  $k_{cat}$  of  $\sim 30 \text{ min}^{-1}$  under optimal assay conditions. An artificial 'RNA ligase' ribozyme has been shown to catalyze the corresponding self-modification reaction with a rate of  $\sim 100 \text{ min}^{-1}$  (Ekland et al., 1995, *Science*, 269, 364). Finally, replacement of a specific residue within the catalytic core of the hammerhead with certain nucleotide analogues gives modified ribozymes that show as much as a 10-fold improvement in catalytic rate (Burgin et al., 1996, *supra*). These findings demonstrate that

ribozymes can promote chemical transformations with catalytic rates that are significantly greater than those displayed *in vitro* by most natural self-cleaving ribozymes. It is then possible that the structures of certain self-cleaving ribozymes may not be optimized to give maximal catalytic activity, or that entirely new RNA motifs could be made that display significantly faster rates for RNA phosphoester cleavage.

An extensive array of site-directed mutagenesis studies have been conducted with the hammerhead to probe relationships between nucleotide sequence and catalytic activity. These systematic studies have made clear that most nucleotides in the conserved core of the hammerhead ribozyme (Forster & Symons, 1987, *Cell*, 49, 211) cannot be mutated without significant loss of catalytic activity. In contrast, a combinatorial strategy that simultaneously screens a large pool of mutagenized ribozymes for RNAs that retain catalytic activity could be used more efficiently to define immutable sequences and to identify new ribozyme variants (Breaker, 1997, *supra*). For example, Joseph and Burke (1993; *J. Biol. Chem.*, 268, 24515) have used an *in vitro* selection approach to rapidly screen for sequence variants of the 'hairpin' self-cleaving RNA that show improved catalytic activity. This approach was successful in identifying two mutations in the hairpin ribozyme that together give a 10-fold improvement in catalytic rate. Although similar *in vitro* selection experiments have been conducted with the hammerhead ribozyme (Nakamaye & Eckstein, 1994, *supra*; Long & Uhlenbeck, 1994, *supra*; Ishizaka et al., 1995, *supra*), none of these efforts have successfully screened full-sized hammerhead ribozymes for all possible combinations

of sequence variants that encompass the entire catalytic core.

Applicant employed *in vitro* selection strategy to comprehensively test whether the natural consensus sequence for the core of the hammerhead ribozyme produces maximal catalytic rates, or whether sequence variants of this ribozyme could catalyze endonuclease reaction similar to or better than the hammerhead ribozyme.

A selection method for self-cleaving ribozymes makes use of the gel-mobility shift that occurs when full-length ribozyme precursors are fragmented and separated by polyacrylamide gel electrophoresis (PAGE). The hammerhead ribozyme can efficiently promote its own cleavage in the presence of the  $Mg^{2+}$ , a metal ion that serves as a cofactor for natural self-cleaving ribozymes and that is also a required component of *in vitro* transcription reactions. As a result, a significant portion of the ribozyme transcripts will self-cleave during preparation by *in vitro* transcription. The ribozyme precursor cannot be isolated from the spurious and unwanted RNA products that are typical of *in vitro* transcription, without losing the portion of precursors that have cleaved during the transcription reaction. Moreover, the best ribozymes present in a mutagenized or random-sequence pool can cleave during transcription, thereby creating a significant impediment to the *in vitro* selection of self-cleaving RNAs. Although ribozymes that cleave during transcription can be recovered by PAGE as part of the *in vitro* selection process, this approach can easily lead to the recovery of smaller 'selfish' RNAs that are not catalytic, but that have an electrophoretic mobility that serendipitously correspond to the cleaved ribozymes (Nakamaye & Eckstein, 1994). In addition, if the selection

process relies on the isolation of ribozymes that cleave during preparation, there is no possibility for selection under alternative reaction conditions that are not compatible with *in vitro* transcription.

5 Example 1: Preparation of intact self-cleaving ribozymes using 'allosteric delay'

Applicant has overcome the problem of ribozyme self-destruction during preparative transcription by making use of 'allosteric ribozyme' constructs that are not cleaved  
10 during transcription, but that remain highly active upon purification. Applicant used an ATP-dependent allosteric ribozyme H3 (Figure 1B; Tang and Breaker, 1997, submitted for publication) in which a hammerhead ribozyme was joined to an RNA aptamer (Sassanfar & Szostak, 1993, Nature 364,  
15 550) that binds adenosine or any of its 5'-phosphorylated derivatives. H3 cleaves separate substrate molecules with a catalytic rate that is highly-dependent on the presence of adenosine or ATP. This conjoined aptamer/ribozyme construct experiences a ~170-fold reduction in catalytic  
20 activity upon addition of 1 mM nucleoside or nucleotide effector.

One of the ribozyme constructs designed by the applicant (Figure 2A; AD-H1) was designed to be analogous to H3, except that the ribozyme and substrate domains in  
25 the new construct are contained within a single molecule. Preparation of AD-H1 in a 3-hr *in vitro* transcription reaction resulted in almost complete inhibition of ribozyme function and, as a result, near complete preservation of the unimolecular ribozyme precursor. All  
30 transcription reactions conducted in this study initially contained 2 mM of each of the four ribonucleoside 5'-triphosphates (NTPs). Therefore, the ATP concentration during transcription far exceeds the  $K_D$  of the ATP-specific

aptamer (10  $\mu$ M), and also exceeds the ATP concentration that was needed to give maximal inhibition of the allosteric ribozyme H3. The purified AD-H1 ribozyme, however, is highly active when allowed to react in the  
5 absence of ATP. AD-H1 is equally active when incubated in 50 mM Tris-HCl (pH 7.5 at 23°C) and 20 mM MgCl<sub>2</sub> (buffer B) or in the transcription buffer (buffer A) in the absence of NTPs. The catalytic rate for AD-H1 (Table 1) is ~3-fold slower than the rate for a similar hammerhead ribozyme  
10 without the appended aptamer domain. However, the catalytic activity of AD-H1 is significantly reduced in the presence of 1 mM ATP or when incubated with the same concentration of NTPs used for *in vitro* transcription. As  
15 a result, the timing of allosteric ribozyme function can be delayed in the presence of specific allosteric effector molecules, thereby allowing the preparation of intact self-cleaving ribozymes by *in vitro* transcription.

In vitro selection of self-cleaving RNAs from the rAD-H2 pool

20 Applicant has made use of this 'allosteric delay' strategy in its efforts to probe the catalytic fitness of the hammerhead ribozyme using *in vitro* selection. The selection was begun with an initial RNA pool (Generation 0; G0), termed 'rAD-H2' (Figure 2C), in which 14  
25 nucleotides of the catalytic core were made random. The randomized region of rAD-H2 is divided into two domains that Applicant identified as 'N<sub>9</sub>' and 'N<sub>5</sub>'. The G0 RNA pool contained  $3 \times 10^{13}$  molecules, corresponding to an average representation of ~100,000 copies for each possible RNA  
30 sequence variant. The rAD-H2 pool was subjected to six successive rounds of *in vitro* selection (see the scheme in Figure 3), which proceeded by first isolating *in vitro* transcribed RNA precursors by PAGE, then by incubating the

RNA pool in the absence of ATP and recovering cleaved RNAs by a gel-mobility shift protocol. Robust self-cleaving activity of the RNA pool was detected after four rounds of selection (Generation 4;G4), with some improvement in the catalytic activity observed with the RNA pool from G6 (Figure 4).

By "randomized region" is meant a region of completely random sequence and/or partially random sequence. By completely random sequence is meant a sequence wherein theoretically there is equal representation of A, T, G and C nucleotides or modified derivatives thereof, at each position in the sequence. By partially random sequence is meant a sequence wherein there is an unequal representation of A, T, G and C nucleotides or modified derivatives thereof, at each position in the sequence. A partially random sequence can therefore have one or more positions of complete randomness and one or more positions with defined nucleotides

Applicant found that 15 of the 38 clones examined from the G6 pool matched the sequence of AD-H2, while 15 additional clones carried a single U to C change at position 21 (Figure 2B; corresponding to position 7 of the hammerhead core in Figure 1A). The identity of this nucleotide is variable in natural hammerhead isolates, and ribozymes with mutations at this position are known to be active in vitro. The  $k_{obs}$  values for AD-H2 with U or C were  $0.068 \text{ min}^{-1}$  and  $0.041 \text{ min}^{-1}$ , respectively, which are consistent with the  $k_{obs}$  of  $0.047 \text{ min}^{-1}$  determined for the ensemble of ribozymes that comprise the G6 pool (Table 1).

Further characterization of the dominant ribozymes reveals that the C and U variants of AD-H2 experience a ~100-fold reduction in catalytic rate in the presence of

1 mM ATP (Table 1). Although AD-H2 experiences dramatic allosteric inhibition, the G4 and G6 RNA populations display considerable activity despite the presence of 1 mM ATP (Figure 4). This finding suggests that a significant  
5 portion of the selected RNAs do not conform to the allosteric ribozyme design and are not inhibited by ATP. Indeed, N<sub>9</sub> and N<sub>5</sub> domains of the remaining eight clones from G6 do not match the hammerhead consensus sequence, but can be grouped into two new classes of Mg<sup>2+</sup>-dependent  
10 self-cleaving ribozymes (Figure 5). Individuals from class I and an individual from class II display *k*<sub>obs</sub> values that match those of the winning hammerhead ribozymes in the absence of ATP, but show no inhibition (class I) or only slight inhibition (class II) in the presence of 1 mM ATP  
15 (Table 1).

Although ribozymes that carry the natural hammerhead consensus sequence dominate the selection that used 20 mM Mg<sup>2+</sup> as cofactor, Applicant wanted to assess the catalytic fitness of the ribozymes under conditions that simulate  
20 the ionic strength and Mg<sup>2+</sup> concentrations of cells (buffer C; 50 mM Tris-HCl (pH 7.5 at 23°C), 250 mM KCl and 2 mM Mg<sup>2+</sup>). To this end, G5 RNA pool was used to carry out additional rounds of selection using buffer C for the ribozyme selection reaction. The population that was  
25 isolated after four rounds using reduced Mg<sup>2+</sup> concentrations (G9-low) displays a rate for self-cleavage of 0.012 min<sup>-1</sup> in buffer C. Sequence analysis of 8 full-length clones from the G9-low pool revealed 6 sequences that conform to class I ribozyme variants (Figure 5A),  
30 while the hammerhead and class II (Figure 5B) variant ribozymes were represented by only one individual each. Dominance by class I ribozymes in the low-concentration Mg<sup>2+</sup> selection is consistent with the fact that these

ribozymes show 2- to 5-fold higher rates in buffer C compared to the AD-H2 ribozyme variants.

#### Characterization of class I variant ribozymes

The two variants of the AD-H2 ribozyme dominate the G6 RNA pool, accounting for nearly 80% of the RNA population, while class I variants represent less than 20% of the population. In order to more efficiently accumulate class I ribozyme sequences for further analysis, we conducted a single additional round of selection (G7-ATP) that exploited the difference in catalytic activity between the dominant hammerhead sequences and the less common class I variants when incubated in the presence of ATP. G6 RNA was incubated for 10 min in the presence of 1 mM ATP under otherwise identical selection conditions. Class I ribozymes are expected to dominate under these new selection conditions because they show no catalytic inhibition in the presence of ATP. As before, the ribozyme's 5'-cleavage product was isolated by PAGE and used to produce the 'G7-ATP' RNA pool. Cloning and sequencing of this pool revealed that nearly 70% of the RNA pool consists of class I ribozymes, while the representation of hammerhead ribozymes dropped to ~25%. The additional class I sequences obtained from G7-ATP are new variants, except for a single repetition of the class I ribozyme v6 (Figure 5A).

The class I ribozyme v2 carries all the most frequently occurring nucleotides in both the N<sub>9</sub> and N<sub>5</sub> domains (Figure 6). Applicant further examined the class I v2 ribozyme by creating the bimolecular ribozyme arrangements termed 'v2 trans' (Figure 7). The v2 trans ribozyme displays catalytic activity. In addition, the new ribozyme cleaves the corresponding RNA substrate at the internucleotide linkage that immediately precedes the



site cleaved by the hammerhead ribozyme. The v2 trans ribozyme cleaves this new site with a  $k_{\text{obs}}$  of  $0.02 \text{ min}^{-1}$  using  $1 \mu\text{M}$  enzyme and trace amounts of substrate in buffer B.

## 5 Ribozyme Engineering

Sequence, chemical and structural variants of Class I and Class II ribozymes can be engineered using the techniques shown above and known in the art to cleave a separate target RNA or DNA in trans. For example, the size of class I and class II ribozymes can, be reduced or increased using the techniques known in the art (Zaug et al., 1986, *Nature*, 324, 429; Ruffner et al., 1990, *Biochem.*, 29, 10695; Beaudry et al., 1990, *Biochem.*, 29, 6534; McCall et al., 1992, *Proc. Natl. Acad. Sci., USA.*, 89, 5710; Long et al., 1994, *Supra*; Hendry et al., 1994, *BBA* 1219, 405; Benseler et al., 1993, *JACS*, 115, 8483; Thompson et al., 1996, *Nucl. Acids Res.*, 24, 4401; Michels et al., 1995, *Biochem.*, 34, 2965; Been et al., 1992, *Biochem.*, 31, 11843; Guo et al., 1995, *EMBO. J.*, 14, 368; Pan et al., 1994, *Biochem.*, 33, 9561; Cech, 1992, *Curr. Op. Struc. Bio.*, 2, 605; Sugiyama et al., 1996, *FEBS Lett.*, 392, 215; Beigelman et al., 1994, *Bioorg. Med. Chem.*, 4, 1715; all are incorporated in its totality by reference herein) For example the aptamer domain (e.g., ATP aptamer) of the ribozymes of either Class I or Class II may not be essential for catalytic activity and hence could be systematically reduced in size using a variety of methods known in the art, to the extent that the overall catalytic activity of the ribozyme is not significantly decreased.

Further rounds of *in vitro* selection strategies described herein and variations thereof can be readily used by a person skilled in the art to evolve additional

nucleic acid catalysts and such new catalysts are within the scope of the instant invention.

#### New self-cleaving ribozymes

The rAD-H2 construct was designed to allow the comprehensive screening of all possible sequence variants of the hammerhead catalytic core. Therefore, the allosteric delay strategy should give a distinct selective advantage to those ribozyme variants that remain active, yet benefit from allosteric inhibition during transcription. Interestingly, two new classes of self-cleaving ribozymes have emerged from the selection that are as active as AD-H2, but that do not undergo ATP-specific allosteric delay. This difference in allosteric inhibition was subsequently used to selectively enrich the RNA pool for class I ribozymes. Specifically, class I ribozymes represent less than 20% of the individual ribozymes that were recovered from the G6 RNA pool. Allosteric inhibition of the dominant hammerhead ribozymes by ATP during the preparation of the G7-ATP RNA pool created a drastic change in the composition of the ribozyme population. The G7-ATP pool consists of ~75% class I variants, thereby making further sequence analysis of this class proceed more efficiently.

Both class I and class II ribozymes have acquired additional mutations outside the N<sub>9</sub> and N<sub>5</sub> randomized domains. These position may be mutation 'hot spots', or may be infrequent but essential mutations that offer a significant selective advantage for those RNAs that have acquired them. Applicant has isolated 18 class I ribozyme variants that have considerable sequence variability, but that carry only a single mutation outside of the randomized N<sub>9</sub> and N<sub>5</sub> domains (Figure 5A). It is likely that most of these class I variants were individually repre-

sented in the original RNA pool and that each variant independently acquired the G to C mutation at position 28 of the aptamer during the *in vitro* selection process. None of the hammerhead ribozymes isolated from G6 carry aptamer  
5 mutations, suggesting that the mutations observed in the new ribozyme classes might be necessary for efficient catalytic function, and that these mutations may occur infrequently.

The class I ribozymes can be divided into separate  
10 ribozyme and substrate domains to create a functional bimolecular complex. This ribozyme presumably interacts with the substrate domain by forming base-paired regions that are analogous to helices I and II of the hammerhead ribozyme (Figure 1). Likewise, the substrate specificity  
15 of class I ribozymes can presumably be altered by changing the sequences of the substrate-binding arms to complement the sequence of the desired substrate molecule, as was achieved with the ribozyme v2 trans (Figure 7). Although the hammerhead and class I ribozymes cleave at different  
20 internucleotide linkages (Figure 1), both ribozymes appear to proceed by a similar chemical mechanism. The hammerhead ribozyme is known to produce a 2',3'-cyclic phosphate at the terminus of the 5'-cleavage product, thereby leaving a 5'-hydroxyl terminus on the 3'-cleavage fragment. The  
25 5'-cleavage products from both the hammerhead- and v2 trans-cleaved substrate RNAs migrate with the major products of alkaline-promoted substrate degradation that are terminated with 2',3'-cyclic phosphates as opposed to the minor products that carry a 2'(3')-phosphate. This is  
30 consistent with the possibility that, like the hammerhead, class I ribozymes utilize a transesterification mechanism to cleave the target RNA phosphodiester linkage. The similarity between the class I ribozymes and the hammerhead appear to end here, since removal of the

aptamer domain results in inactivation of the v2 ribozyme. Therefore, at least a portion of the mutant aptamer domain appears to be required for class I catalytic function.

The natural consensus sequence for the hammerhead catalytic core also was not improved upon by *in vitro* selection using a low  $Mg^{2+}$  concentration (buffer C). Here, the class I variant ribozymes come to dominate the G9-low RNA pool, however, the catalytic advantage of class I ribozymes compared to the hammerhead ribozymes that were isolated at G6 is subtle (Table 1). Although the class II ribozyme v1 displays catalytic rates that are similar to both the hammerhead and class I ribozymes in buffer B, and to class I ribozymes in buffer C (Table 1). However, this ribozyme or related variants do not comprise a major portion of either the G6 or the G9-low RNA pools. If the mutations acquired by the two representatives of class II are necessary for efficient catalytic function, then these ribozymes would have been present at a much lower frequency than either the hammerhead or class I ribozymes. As a result, this class of catalyst would not be expected to dominate the selected RNA pools due to their infrequent occurrence in the G0 and subsequent RNA pools.

Applicant has used ATP-controlled allosteric ribozymes to create self-cleaving hammerhead ribozymes that can be isolated as intact precursors from *in vitro* transcription reactions. Hammerhead ribozyme, as well as other ribozymes, could be designed that can be controlled by different allosteric effector molecules by judicious coupling of distinct aptamer and ribozyme domains. In the instant study, the ATP-dependent allosteric ribozyme design has been particularly beneficial for preparing unimolecular hammerhead ribozymes for use in kinetic analyses. Applicant has employed this concept of

allosteric delay in the design of a randomized RNA construct that was used to probe the catalytic fitness of the hammerhead ribozyme.

Using the allosteric delay strategy, Applicant found  
5 that the dominant class of ribozymes to emerge after six rounds of *in vitro* selection included sequences that matched the natural consensus of the catalytic core, including the re-identification of the variable-nucleotide site at position 7. This comprehensive screening of all  
10 possible sequence variations within the 14-nucleotide core region is expected to have simultaneously tested the catalytic fitness of nearly 270 million RNA molecules.

Finally, although the allosteric selection construct was heavily biased in favor of the hammerhead ribozyme,  
15 the selection produced two new ribozymes that cleave RNA with rates that are comparable to the allosteric hammerhead ribozymes. This finding, taken together with other reports of the *in vitro* selection of unique self-cleaving ribozymes, suggests that many novel ribozymes  
20 with similar catalytic activities have yet to be found. The application of *in vitro* selection using an unbiased starting pool of random-sequence RNA is likely to produce novel self-cleaving ribozymes with catalytic rates that meet or perhaps even exceed those of natural ribozymes.

## 25 Materials and Methods

### Synthetic oligonucleotides and transcription-template constructs

Synthetic DNA and RNA oligonucleotides were prepared  
(Keck Biotechnology Resource Laboratory, Yale University)  
30 by standard solid-phase methods, purified by denaturing PAGE (8 M urea, 89 mM Tris-borate, 2 mM EDTA), and isolated by crush-soaking in 10 mM Tris-HCl (pH 7.5 at 23°C), 200 mM NaCl and 1 mM EDTA. The 2'-TBDMS group of the

synthetic RNA substrate (5'-GCCGUAGGUUGCCC) was removed by 24-hr treatment with triethylamine trihydrofluoride (15  $\mu$ l per AU<sub>260</sub> crude RNA). Purified RNA substrate was [5'-<sup>32</sup>P]-labeled with T4 polynucleotide kinase and [ $\gamma$ -<sup>32</sup>P]-ATP, then  
5 repurified by PAGE.

#### In vitro transcription

For each RNA sequence or RNA pool, 10 to 100 pmoles of DNA template was transcribed in buffer A (50 mM Tris-HCl (pH 7.5 at 23°C), 15 mM MgCl<sub>2</sub>, 5 mM dithiothreitol and  
10 2 mM spermidine) containing 2 mM each of the four ribonucleoside triphosphates. RNA synthesis was initiated by the addition of T7 RNA polymerase to a final concentration of 12 U  $\mu$ L<sup>-1</sup> and incubated for 1 to 3 hours at 37°C. Internally-labeled transcripts were prepared by  
15 the inclusion of 20  $\mu$ Ci of [ $\alpha$ -<sup>32</sup>P]-UTP in the transcription reactions. Transcription products were separated by denaturing 10% PAGE, visualized by autoradiography or by electronic imaging (PhosphorImager, Molecular Dynamics) and the ribozymes were recovered from excised gel  
20 particles by crush-soaking. Concentrations of purified RNAs were established by liquid scintillation counting.

#### In vitro selection

G0 pool RNA was prepared by transcribing 100 pmoles of double-stranded DNA in a 100- $\mu$ L reaction volume,  
25 corresponding to a population of rAD-H2 templates that is expected to include all possible sequence combinations within the catalytic core of the hammerhead ribozyme. This template DNA was prepared by extending 200 pmoles primer 2 (5'-GAATTCTAATAC-GACTCACTATAGGAAGAGATGGCGAC) in the  
30 presence of 200 pmoles of the oligonucleotide 5'-TTTGAGGCGACCTACCACTCTCGTGG(N)<sub>5</sub>TTGCTGCGACCGAAGTCGCACAGTTTC-TTCCCAA(N)<sub>9</sub>GTCGCCATCTCTTCC (where N indicates an equal mix

of the four nucleotides) with Taq polymerase under polymerase chain reaction (PCR) conditions. The PCR extension reaction was conducted in a total of 100  $\mu\text{L}$  containing 0.05 U  $\mu\text{L}^{-1}$  Taq polymerase, 50 mM KCl, 1.5 mM MgCl<sub>2</sub>, 10 mM Tris-HCl (pH 8.3 at 23°C), 0.01 % gelatin, and 0.2 mM each dNTP for 1 cycle of 20 sec at 92°C, 20 sec at 50° C and 30 sec at 72°C.

For the initial round of selection, a total of 50 pmoles rAD-H2 RNA was incubated in a 50  $\mu\text{L}$  reaction mixture containing 50 mM Tris-HCl (pH 7.5 at 23°C) and 20 mM MgCl<sub>2</sub> (buffer B) for 1 hr at room temperature. The reaction was terminated by the addition of an equal volume of PAGE loading buffer (8 M urea, 5 mM Tris-borate (pH 8.3 at 23°C), 0.3 M sucrose, 50 mM Na<sub>2</sub>EDTA, 0.02% w/v xylene cyanol, and 0.02% w/v bromophenyl blue) and the products were separated by denaturing 10% PAGE. Reaction products were visualized by autoradiography, phosphorimage, or by UV shadowing. Great care was taken to optimize the separation between uncleaved precursor RNAs and the 5' cleavage fragments. The cleaved ribozymes were isolated by crush/soaking the corresponding gel fragments and were concentrated by precipitation with ethanol. Recovered RNAs were reverse transcribed using 30 pmoles of primer 1 (5'-TTTGATGGCGACCTACCACTCTC-GTGG) in a 25  $\mu\text{L}$  reaction containing 10 U  $\mu\text{L}^{-1}$  SuperScript reverse transcriptase (BRL) and incubated at 37°C for 30 min in the buffer supplied by the manufacturer. The resulting DNA was amplified by PCR as described above for 25 cycles. This entire process was repeated for addition selection rounds with some alterations. Specifically, transcription of subsequent generations was conducted using ~12 pmoles of PCR DNA from the preceding generation in a total of 50  $\mu\text{L}$  as described above. RNA populations G3-G6 were twice

purified by PAGE to eliminate deletion mutants. After G3, only 50% of the cDNA from the selected RNA molecules was used for amplification by PCR. Finally, the reaction time for the sixth round of selection was reduced to 3 min.

5       The 'G7-ATP' pool was generated by selecting RNAs from the G6 pool that are active in the presence of ATP. This selection proceeded like that of the previous rounds, but the ribozyme selection reaction was conducted for 10 min in buffer B containing 1 mM ATP. The 'G9-low' pool was  
10       derived from G5 RNA, which was subjected to four additional rounds of selection using a ribozyme reaction buffer containing 50 mM Tris-HCl (pH 7.5 at 23°C), 250 mM KCl, and 2 mM MgCl<sub>2</sub> (buffer C). The low magnesium selection reactions were carried out at 23°C for 1 hr, or 10 min for  
15       the final round. Individuals from all populations of interest were analyzed by cloning (TA cloning kit, Invitrogen) and sequencing (ThermalSequenase Kit, Amersham).

#### Ribozyme assays

20       Self-cleaving ribozyme assays were conducted with internally-labeled precursor RNAs in buffers A, B or C as indicated for each experiment. Bimolecular ribozyme assays were conducted with trace amounts of [5'-<sup>32</sup>P]-labeled substrate RNA and 1 μM of internally-labeled RNA enzyme.  
25       The results were analyzed by denaturing 10% PAGE and were visualized and analyzed by autoradiography or by PhosphorImager (Molecular Dynamics). For kinetic assays, a series of time points were made that best represented the initial rate of ribozyme cleavage. Catalytic rates  
30       (*k*<sub>obs</sub>) were obtained by plotting the fraction of substrate cleaved versus time and establishing the slope of the line that represents the initial velocity of the reaction. For assays conducted in 1 mM ATP, the precursor RNA was



preincubated with ATP for 10 min in the absence of  $Mg^{2+}$ . This eliminated the burst kinetics that are observed with reactions that are initiated by the simultaneous addition of ATP and  $Mg^{2+}$ . Replicate experiments gave  $k_{obs}$  values  
5 that differed by less than 20% and the values reported are averages of two or more repetitions.

Cleavage sites for H1 and v2 trans ribozymes were determined by PAGE separation of ribozyme reactions using a partial alkaline digest of the synthetic substrate as a  
10 marker. The partial alkaline digest was made by incubation of a trace amount of  $[5'-^{32}P]$ -labeled substrate RNA in 100 mM NaOH for 5 min.

#### Diagnostic uses

Enzymatic nucleic acids of this invention may be used  
15 as diagnostic tools to examine genetic drift and mutations within diseased cells or to detect the presence of target RNA in a cell. The close relationship between ribozyme activity and the structure of the target RNA allows the detection of mutations in any region of the molecule which  
20 alters the base-pairing and three-dimensional structure of the target RNA. By using multiple ribozymes described in this invention, one may map nucleotide changes which are important to RNA structure and function *in vitro*, as well as in cells and tissues. Cleavage of target RNAs with  
25 ribozymes may be used to inhibit gene expression and define the role (essentially) of specified gene products in the progression of disease. In this manner, other genetic targets may be defined as important mediators of the disease. These experiments will lead to better  
30 treatment of the disease progression by affording the possibility of combinational therapies (e.g., multiple ribozymes targeted to different genes, ribozymes coupled with known small molecule inhibitors, or intermittent

treatment with combinations of ribozymes and/or other chemical or biological molecules). Other in vitro uses of ribozymes of this invention are well known in the art, and include detection of the presence of mRNAs associated with  
5 disease condition. Such RNA is detected by determining the presence of a cleavage product after treatment with a ribozyme using standard methodology.

In a specific example, ribozymes which can cleave only wild-type or mutant forms of the target RNA are used  
10 for the assay. The first ribozyme is used to identify wild-type RNA present in the sample and the second ribozyme will be used to identify mutant RNA in the sample. As reaction controls, synthetic substrates of both wild-type and mutant RNA will be cleaved by both  
15 ribozymes to demonstrate the relative ribozyme efficiencies in the reactions and the absence of cleavage of the "non-targeted" RNA species. The cleavage products from the synthetic substrates will also serve to generate size markers for the analysis of wild-type and mutant RNAs  
20 in the sample population. Thus each analysis will require two ribozymes, two substrates and one unknown sample which will be combined into six reactions. The presence of cleavage products will be determined using an RNase protection assay so that full-length and cleavage  
25 fragments of each RNA can be analyzed in one lane of a polyacrylamide gel. It is not absolutely required to quantify the results to gain insight into the expression of mutant RNAs and putative risk of the desired phenotypic changes in target cells. The expression of mRNA whose  
30 protein product is implicated in the development of the phenotype is adequate to establish risk. If probes of comparable specific activity are used for both transcripts, then a qualitative comparison of RNA levels will be adequate and will decrease the cost of the initial

diagnosis. Higher mutant form to wild-type ratios will be correlated with higher risk whether RNA levels are compared qualitatively or quantitatively.

#### Additional Uses

- 5 Potential usefulness of sequence-specific enzymatic nucleic acid molecules of the instant invention might have many of the same applications for the study of RNA that DNA restriction endonucleases have for the study of DNA (Nathans et al., 1975 *Ann. Rev. Biochem.* 44:273). For  
10 example, the pattern of restriction fragments could be used to establish sequence relationships between two related RNAs, and large RNAs could be specifically cleaved to fragments of a size more useful for study. The ability to engineer sequence specificity of the ribozyme is ideal  
15 for cleavage of RNAs of unknown sequence.

Other embodiments are within the following claims.

Table I: Cleavage Kinetics of Allosteric Ribozymes

Construct	$k_{obs}$ (min <sup>-1</sup> )			ATP Mg <sup>2+</sup> (mM)
	-	+	-	
	20	20	2	
AD-H1	0.18	-	0.011	
G6 pool	0.047	-	-	
AD-H2 (U)	0.068	0.00051	0.0051	
AD-H2 (C)	0.041	0.00050	0.0021	
class I (v1)	0.055	0.043	0.0032	
class I (v2)	0.070	0.053	0.011	
class II (v1)	0.054	0.016	0.012	

Table II

Table II: 2.5  $\mu$ mol RNA Synthesis Cycle

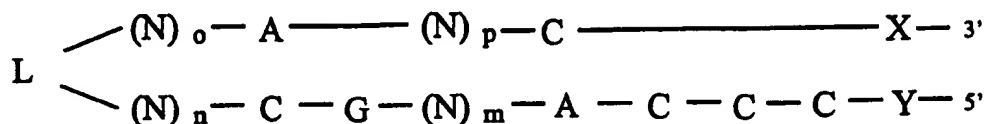
Reagent	Equivalents	Amount	Wait Time*
Phosphoramidites	6.5	163 $\mu$ L	2.5
S-Ethyl Tetrazole	23.8	238 $\mu$ L	2.5
Acetic Anhydride	100	233 $\mu$ L	5 sec
N-Methyl Imidazole	186	233 $\mu$ L	5 sec
TCA	83.2	1.73 mL	21 sec
Iodine	8.0	1.18 mL	45 sec
Acetonitrile	NA	6.67 mL	NA

\* Wait time does not include contact time during delivery.

Claims

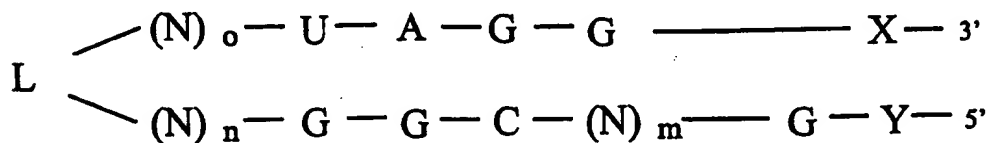
1. A nucleic acid molecule with an endonuclease activity having the formula I:

5



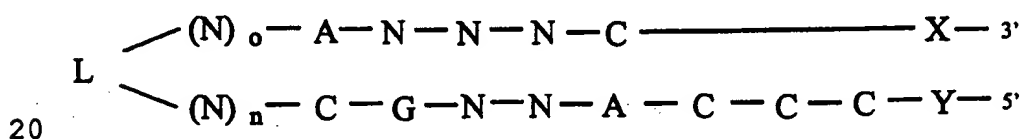
wherein, each N represents independently a nucleotide or a non-nucleotide linker, which may be same or different;  
 10 X and Y are independent oligonucleotides of length sufficient to stably interact with a target nucleic acid molecule; o and n are integers greater than or equal to 1, wherein if (N)<sub>o</sub> and (N)<sub>n</sub> are nucleotides, (N)<sub>o</sub> and (N)<sub>n</sub> may interact by hydrogen bond interaction; p and m are  
 15 selected independently from the integers 0, 1, 2, 3, 4 or 5; L is a linker which may be present or absent, wherein said linker, when present, is a nucleotide and/or a non-nucleotide linker, wherein said nucleotide linker comprises a single-stranded and/or double-stranded region;  
 20 \_\_\_\_\_ represents a chemical linkage; and A, C and G represent adenosine, cytidine and guanosine nucleotides, respectively.

2. A nucleic acid molecule with catalytic activity  
 25 having the formula II:



wherein, each N represents independently a nucleotide or a non-nucleotide linker, which may be same or different; X and Y are independent oligonucleotides of length sufficient to stably interact with a target nucleic acid molecule; o and n are integers greater than or equal to 1, wherein if (N)<sub>o</sub> and (N)<sub>n</sub> are nucleotides, (N)<sub>o</sub> and (N)<sub>n</sub> may interact by hydrogen bond interaction; m is selected independently from the integers 0, 1, 2, 3, 4 or 5; L is a linker which may be present or absent, wherein said linker, when present, is a nucleotide and/or a non-nucleotide linker, wherein said nucleotide linker comprises a single-stranded and/or double-stranded region; \_\_\_\_\_ represents a chemical linkage; and A, C, U and G represent adenosine, cytidine, uridine and guanosine nucleotides, respectively.

3. A nucleic acid molecule with catalytic activity having the formula III:

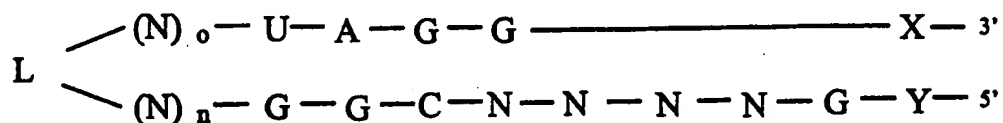


wherein, each N represents independently a nucleotide or a non-nucleotide linker, which may be same or different; X and Y are independent oligonucleotides of length sufficient to stably interact with a target nucleic acid molecule; o and n are integers greater than or equal to 1, wherein if (N)<sub>o</sub> and (N)<sub>n</sub> are nucleotides, (N)<sub>o</sub> and (N)<sub>n</sub> may interact by hydrogen bond interaction; L is a linker which may be present or absent, wherein said linker, when present, is a nucleotide and/or a non-nucleotide linker, wherein said nucleotide linker comprises a single-stranded and/or double-stranded region; \_\_\_\_\_ represents a

46

chemical linkage; and A, C and G represent adenosine, cytidine and guanosine nucleotides, respectively.

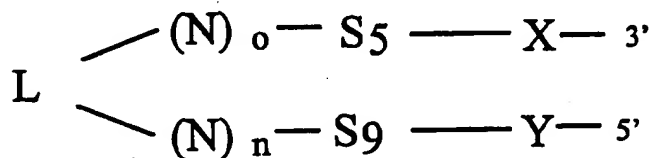
4. A nucleic acid molecule with catalytic activity  
5 having the formula IV:



wherein, each N represents independently a nucleotide or  
10 a non-nucleotide linker, which may be same or different;  
X and Y are independent oligonucleotides of length  
sufficient to stably interact with a target nucleic acid  
molecule; o and n are integers greater than or equal to 1,  
wherein if (N)<sub>o</sub> and (N)<sub>n</sub> are nucleotides, (N)<sub>o</sub> and (N)<sub>n</sub>  
15 may interact by hydrogen bond interaction; L is a linker  
which may be present or absent, wherein said linker, when  
present, is a nucleotide and/or a non-nucleotide linker,  
wherein said nucleotide linker comprises a single-stranded  
and/or double-stranded region; \_\_\_\_\_ represents a  
20 chemical linkage; and A, C, U and G represent adenosine,  
cytidine, uridine and guanosine nucleotides, respectively.

5. A nucleic acid molecule with catalytic activity  
having the formula V:

25



47

wherein, each N represents independently a nucleotide or a non-nucleotide linker, which may be same or different; X and Y are independent oligonucleotides of length sufficient to stably interact with a target nucleic acid molecule; o and n are integers greater than or equal to 1, wherein if (N)<sub>o</sub> and (N)<sub>n</sub> are nucleotides, (N)<sub>o</sub> and (N)<sub>n</sub> may interact by hydrogen bond interaction; L is a linker which may be present or absent, wherein said linker, when present, is a nucleotide and/or a non-nucleotide linker, wherein said nucleotide linker comprises a single-stranded and/or double-stranded region; \_\_\_\_\_ represents a chemical linkage; S<sub>5</sub> is an oligonucleotide containing a sequence selected from the group consisting of 5'-AUGUC-3', 5'-ACGUC-3', 5'-ACGGC-3', 5'-ACCUC-3', 5'-AAGGC-3', 5'-AUGGC-3', 5'-AUGCC-3', 5'-ACUCC-3', 5'-AUGAC-3', 5'-ACGAC-3', 5'-UUAGG-3', and 5'-CUAGG-3'; and S<sub>9</sub> is an oligonucleotide containing a sequence selected from the group consisting of 5'-CCCAGUGCC-3', 5'-CCCAGUGCA-3', 5'-CCCAUGCA-3', 5'-CCCAUGCC-3', 5'-CCCAUGCU-3', 5'-CCCAUAGCA-3', 5'-CCCAACGCA-3', 5'-CCCACCGCA-3', 5'-CCCAGAGCA-3', 5'-CCCAACGCU-3', 5'-CCCAUUGCA-3', 5'-GUAGACGGA-3', and 5'-GGUUUCGGG-3'.

6 The nucleic acid molecules of any of claims 1-5 further comprises a cytidine residue immediately 3' of said (N)<sub>n</sub>.

7. The nucleic acid molecules of any of claims 1-5, wherein said L is nucleotide linker.

30

8. The nucleic acid molecule of claim 7, wherein said nucleotide linker is a nucleic acid aptamer.



9. The nucleic acid molecule of claim 8, wherein said aptamer is an ATP aptamer.

10. The nucleic acid molecule of claim 6, wherein said L is nucleotide linker.

11. The nucleic acid molecule of claim 10, wherein said nucleotide linker is a nucleic acid aptamer.

10 12. The nucleic acid molecule of claim 11, wherein said aptamer is a ATP aptamer.

13. The nucleic acid molecule of any of claims 1-5, wherein said nucleic acid cleaves a separate nucleic acid molecule.

14. The nucleic acid molecule of claim 13, wherein said separate nucleic acid molecule is RNA.

20 15. The nucleic acid molecule of claim 13, wherein said nucleic acid comprises between 12 and 100 bases complementary to said separate nucleic acid molecule.

16. The nucleic acid molecule of claim 13, wherein said nucleic acid comprises between 14 and 24 bases complementary to said separate nucleic acid molecule.

17. A cell including the nucleic acid molecule of any of claims 1-5.

30

18. The cell of claim 17, wherein said cell is a mammalian cell.

19. The cell of claim 18, wherein said cell is a human cell.

20. An expression vector comprising nucleic acid  
5 sequence encoding at least one of the nucleic acid molecule of any of claims 1-5, in a manner which allows expression of that nucleic acid molecule.

21. A cell including the expression vector of claim  
10 20.

22. The cell of claim 21, wherein said cell is a mammalian cell.

15 23. The cell of claim 21, wherein said cell is a human cell.

24. A pharmaceutical composition comprising the nucleic acid molecule of any of claims 1-5.  
20

25. A method for modulating expression of a gene in a plant cell by administering to said cell the nucleic acid molecule of any of claims 1-5.

25 26. A method for modulating expression of gene in a mammalian cell by administering to said cell the nucleic acid molecule of any of claims 1-5.

27. A method of cleaving a separate nucleic acid  
30 comprising, contacting the nucleic acid molecule of any of claims 1-5 with said separate nucleic acid molecule under conditions suitable for the cleavage of said separate nucleic acid molecule.

28. The method of claim 27, wherein said cleavage is carried out in the presence of a divalent cation.

29. The method of claim 28, wherein said divalent  
5 cation is  $Mg^{2+}$ .

30. The nucleic acid molecule of claims 1-5, wherein said nucleic acid is chemically synthesized.

10 31. The expression vector of claim 19, wherein said vector comprises:

- a) a transcription initiation region;
- b) a transcription termination region;
- c) a gene encoding at least one said nucleic acid  
15 molecule; and

wherein said gene is operably linked to said initiation region and said termination region, in a manner which allows expression and/or delivery of said nucleic acid molecule.

20

32. The expression vector of claim 20, wherein said vector comprises:

- a) a transcription initiation region;
- b) a transcription termination region;
- 25 c) an open reading frame;
- d) a gene encoding at least one said nucleic acid molecule, wherein said gene is operably linked to the 3'-end of said open reading frame; and

wherein said gene is operably linked to said  
30 initiation region, said open reading frame and said termination region, in a manner which allows expression and/or delivery of said nucleic acid molecule.

33. The expression vector of claim 20, wherein said vector comprises:

- a) a transcription initiation region;
- b) a transcription termination region;
- 5 c) an intron;
- d) a gene encoding at least one said nucleic acid molecule; and

wherein said gene is operably linked to said initiation region, said intron and said termination region, in  
10 a manner which allows expression and/or delivery of said nucleic acid molecule.

34. The expression vector of claim 20, wherein said vector comprises:

- 15 a) a transcription initiation region;
  - b) a transcription termination region;
  - c) an intron;
  - d) an open reading frame;
  - e) a gene encoding at least one said nucleic acid
- 20 molecule, wherein said gene is operably linked to the 3'-end of said open reading frame; and

wherein said gene is operably linked to said initiation region, said intron, said open reading frame and said termination region, in a manner which allows  
25 expression and/or delivery of said nucleic acid molecule.

35. A method of selecting a nucleic acid with catalytic activity comprising the steps of:

- a) providing a ligand binding nucleic acid region  
30 and a randomized nucleic acid region attached to said ligand binding nucleic acid region; and
- b) selecting for nucleic acid molecules with said catalytic activity in the presence of said ligand.

36. The method of claim 35, wherein said ligand binding region is an ATP aptamer.

5 37. The method of claim 36, wherein said ligand is ATP.

38. A nucleic acid molecule with catalytic activity selected using the method of claim 35.

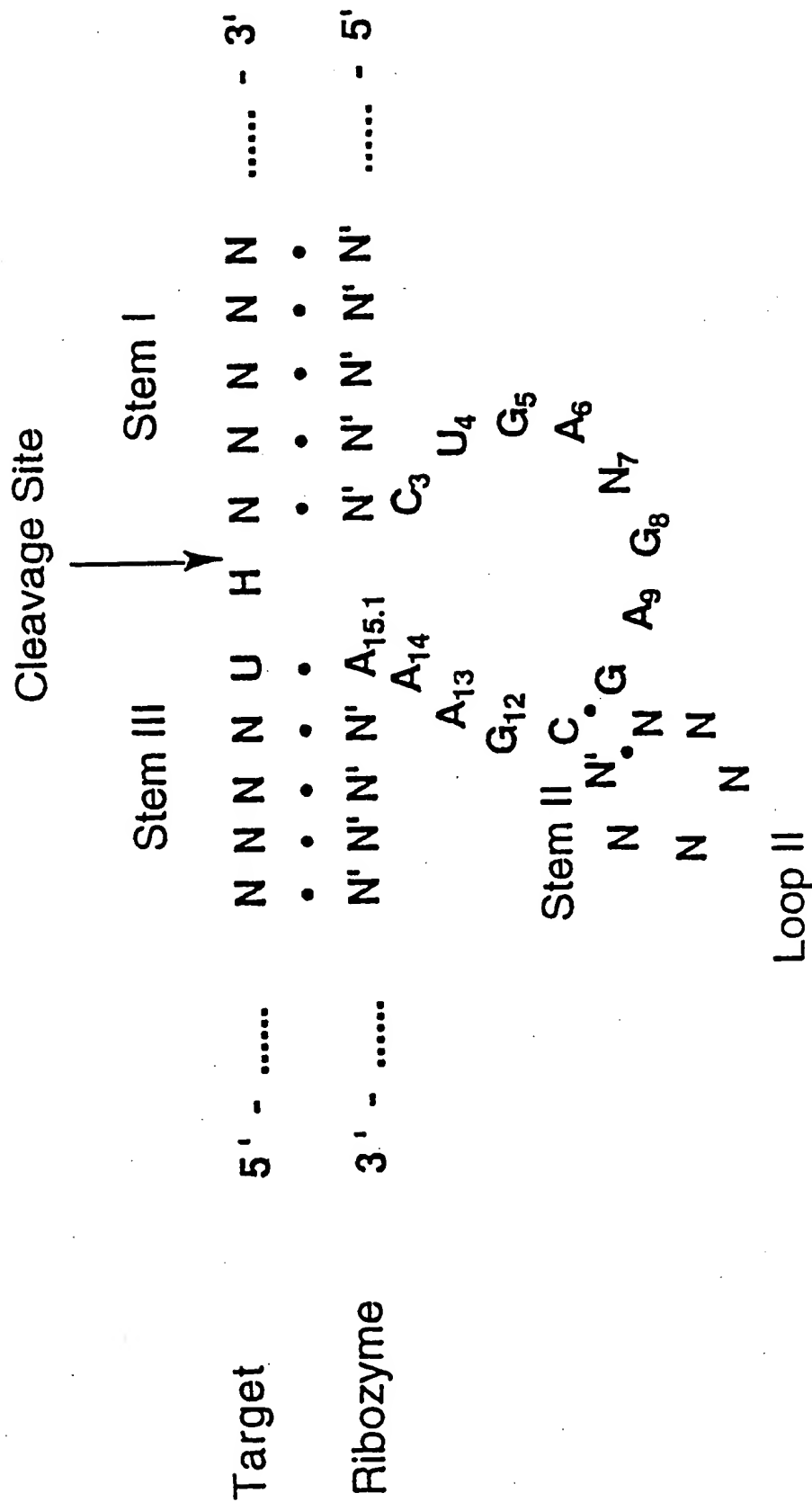
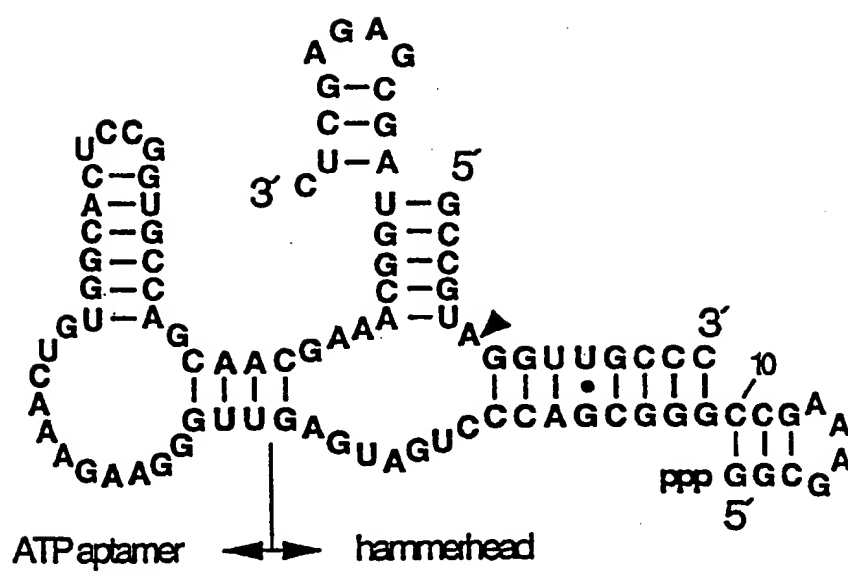


Fig. 1A



### H3

**Fig. 1B**

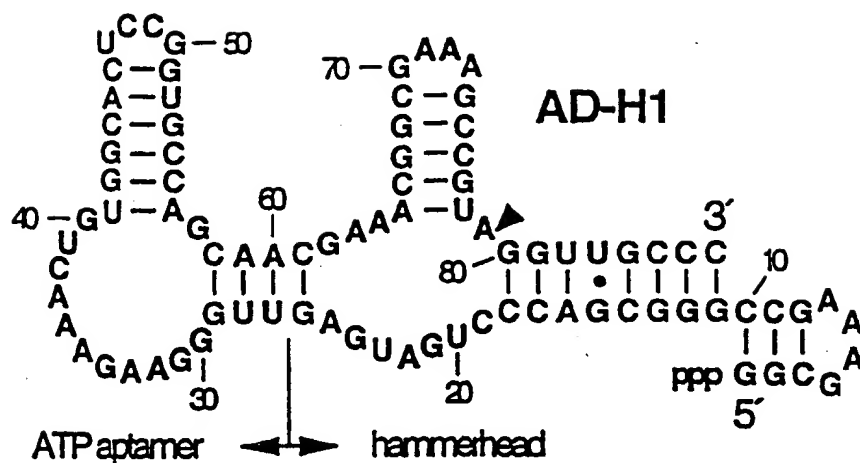


Fig. 2A

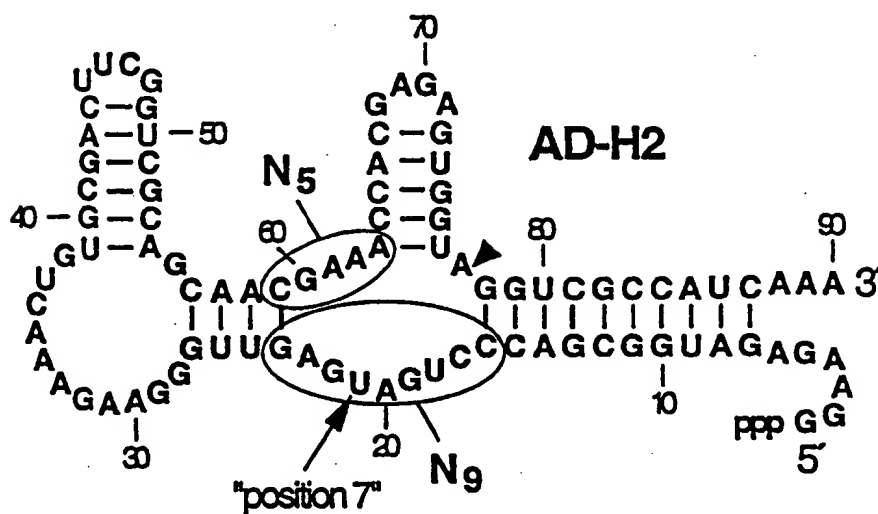


Fig. 2B

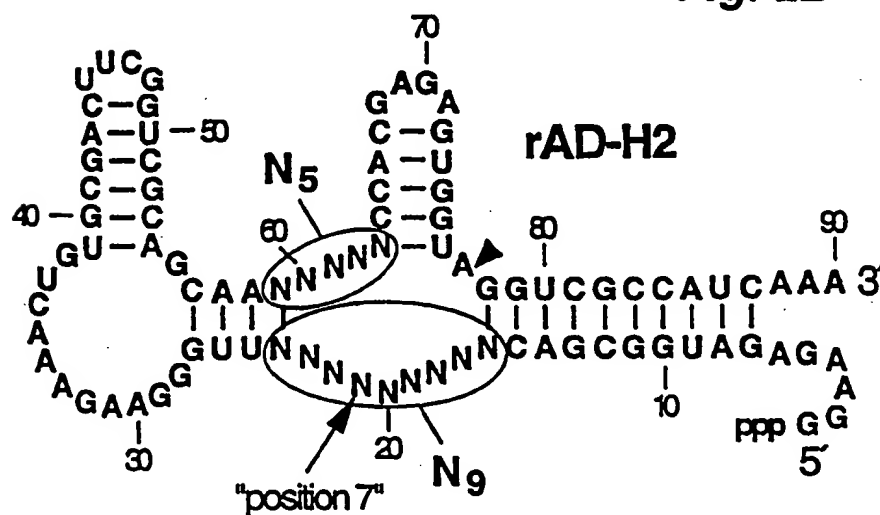


Fig. 2C



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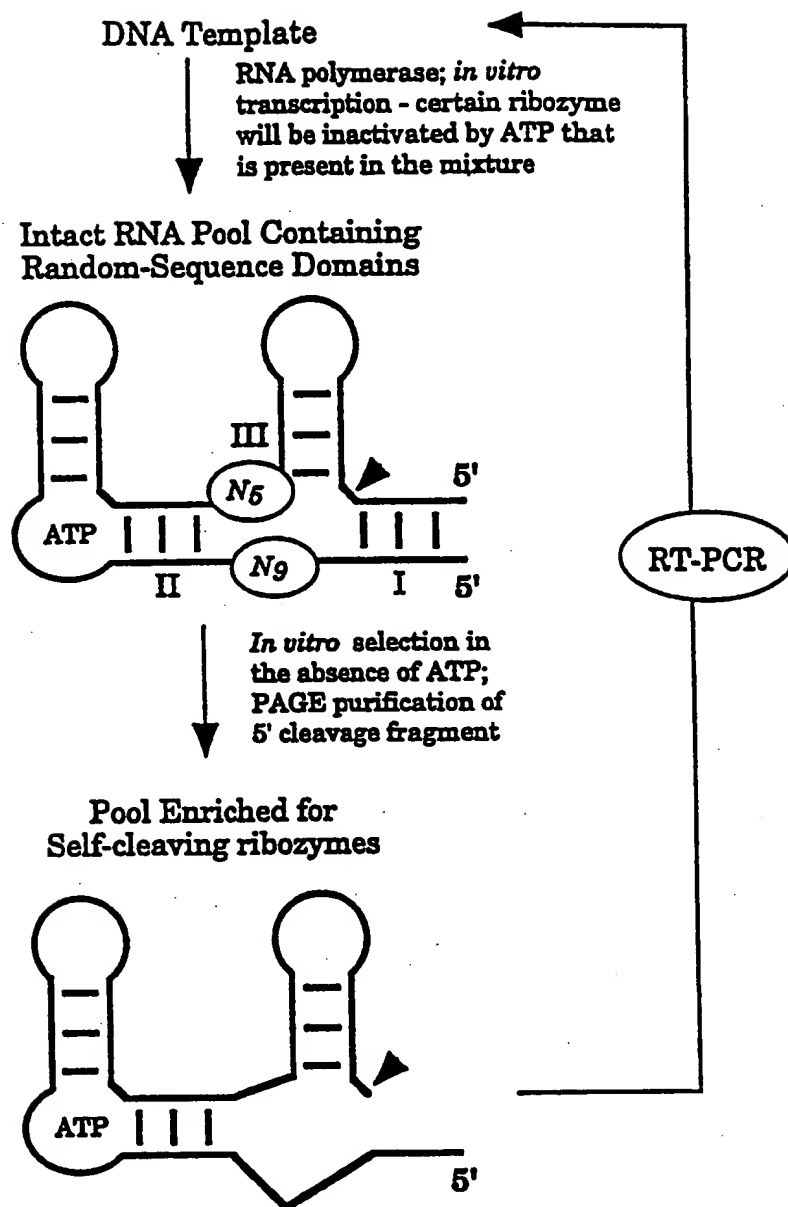
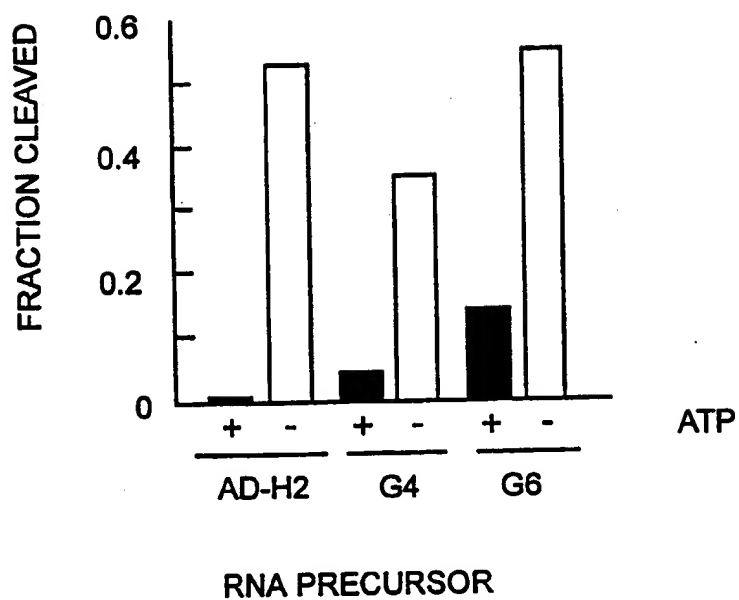


Fig. 3

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**Fig. 4**

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**Class I****Fig. 5A**

	5'	N <sub>9</sub>	N <sub>5</sub>	3'
Generation 6	v1	CCCAGUGCC	AUGUC	
	v2	CCCAGUGCA	ACGUC	
	v3	CCCAGUGCA	ACGGC	
	v4	CCCAAUGCA	ACGUC	
	v5	CCCAAUGCC	ACCUC	
	v6	CCCAGUGCA	AAGGC	
	v7	CCCAAUGCU	AUGGC	
Generation 7-ATP	v6 (1)			
	v8	CCCAGUGCA	AUGCC	
	v9	CCCAUAGCA	AUGGC	
	v10	CCCAAUGCA	ACUCC	
	v11	CCCAACGCA	ACGGC	
	v12	CCCACCGCA	AUGAC	
	v13	CCCACCGCA	AUGGC	
	v14	CCCAAUGCC	AUGGC	
	v15	CCCAGAGCA	ACGAC	
	v16	CCCAACGCU	AUGGC	
	v17	CCCAUUGCA	ACGGC	
Generation 9 Low	v2 (3)			
	v4 (1)			
	v18	CCCAUAGCA	ACGAC	
	v19	CCCAGUGCA	ACGAC	

**Class II****Fig. 5B**

	5'	N <sub>9</sub>	N <sub>5</sub>	3'
v1		GUAGACGGA	UUAGG	
v2		GGUUUCGGG	CUAGG	

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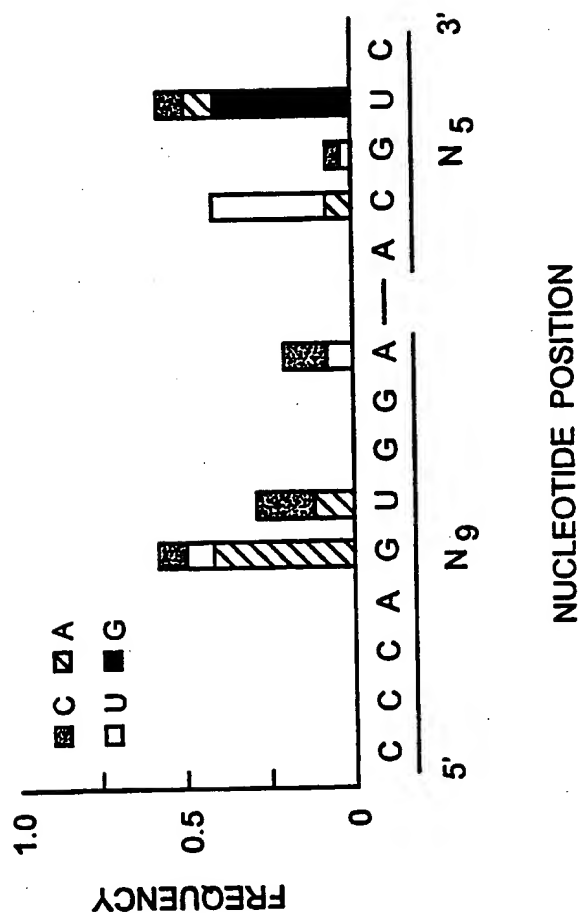


Fig. 6

